From the 11/26/2021 release of VAERS data:

Found 99,943 cases where Vaccine targets COVID-19 (COVID19) and Hospitalized

Government Disclaimer on use of this data

Table

Age	Count	Percent
< 3 Years	29	0.03%
3-6 Years	11	0.01%
6-9 Years	9	0.01%
9-12 Years	16	0.02%
12-17 Years	1,593	1.59%
17-44 Years	16,995	17%
44-65 Years	19,261	19.27%
65-75 Years	11,855	11.86%
75+ Years	15,469	15.48%
Unknown	34,705	34.72%
TOTAL	99,943	100%

Case Details (Sorted by Appearance Date)

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VAERS ID: 902479 (history)
Form: Version 2.0
Age: 46.0

Sex: Female
Location: Kansas
Vaccinated: 2020-12-14
Onset: 2020-12-14
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-15

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Chest pain, Feeling abnormal, Flushing, Intensive care
SMQs:, Anaphylactic reaction (broad), Dementia (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: No
Preexisting Conditions: unknown
Allergies: No
Diagnostic Lab Data: Unknown but sent to SICU for monitoring
CDC Split Type:

Write-up: rPfizer-BionNTech COVID-19 Vaccine EUA 5-7 minutes after the vaccine Associate stated she did not feel right, mentioned chest pain. "My chest feels funny. It feels like when you have really bad heartburn coming on". "I feel flushed like when you get contrast for a CT". Pulse 90 BP 160/90 checked later 130/90

VAERS ID: 903260 (history)
Form: Version 2.0
Age: 60.0
Sex: Female
Location: Alaska
Vaccinated: 2020-12-16
Onset: 2020-12-16
Days after vaccination: 0

Submitted: 0000-00-00
Entered: 2020-12-17

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Condition aggravated, Cough, Tracheomalacia, Wheezing
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Asthma/bronchospasm (broad), Eosinophilic pneumonia (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations: MMR and typhoid - hx of "passing out" and "throat closing"; unsure of vaccination dates, not recent
Other Medications: Acetaminophen, Albuterol, Excedrin Migraine, B complex vitamins tab, Cholecalciferol, Cyclobenzaprine, Duloxetine Dr, Esomeprazole, Norco, Analpram-HC rectal cream, hydroxyzine hydrochloride, ipratropium nasal spray, Duoneb, levothyroxine,
Current Illness:
Preexisting Conditions: asthma, diverticulosis, tracheomalacia, migraines, sciatica, sleep apnea, hiatal hernia, GERD, T2DM
Allergies: Anaphylaxis to amide, halogenated and ester anesthetics, and zofran; history of swelling with bee venom; Hives, shortness of breath and rash with peaches and sulfamethoxazole-trimethoprim; SVT with rizatriptan and sumatriptan
Diagnostic Lab Data:
CDC Split Type:

Write-up: Approximately 10 minutes after vaccine administration, patient reported wheezing and coughing. Patient received epinephrine IM, IV Benadryl, IV solumedrol and racemic epinephrine SVN. Patient never developed a rash, hypotension, swelling of the lips, mouth or tongue, other GI side effects . Per ER attending and admitting physician, this reaction seems to be a clear exacerbation of the patient''s tracheomalacia. The patient was more responsive to racemic epi SVN as opposed to IM epi. Patient admits that psychological stress may have been a component of her symptoms. The admitting physician does not consider this to be an anaphylactic reaction to the vaccination.

VAERS ID: 902854 (history)
Form: Version 2.0
Age: 49.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-08

Onset: 2020-12-08
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-17

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Anxiety, Cough, Fatigue, Hypertension, Limb discomfort, Peripheral circulatory failure, Swollen tongue, Throat irritation, Throat tightness
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypertension (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: VITAMIN B12 [VITAMIN B12 NOS]
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Food allergy
Allergies:
Diagnostic Lab Data: Test Date: 20201208; Test Name: BLOOD PRESSURE SYSTOLIC; Result Unstructured Data: Test Result:175 mmHg; Test Date: 20201208; Test Name: HEART RATE; Result Unstructured Data: Test Result:110; Comments: bpm; Test Date: 20201208; Test Name: PULSE OXIMETRY; Result Unstructured Data: Test Result:NO TRACE
CDC Split Type: GBPFIZER INC2020486806

Write-up: Anaphylaxis; throat closing; tongue swelling; Peripheral shutdown; This is a spontaneous report from a contactable physician and pharmacist received from the Agency. The regulatory authority report number is GB-MHRA-WEBCOVID-20201209083957 and ADR 24541453-001 and ADR 24541453-002. A 49-year-old female patient (healthcare worker) received BNT162b2 vaccine (Batch/lot: EJ0553) on 08Dec2020, via an unspecified route of administration route at single dose for COVID-19 immunization.The patient had a pertinent medical history of food allergy (Lemon/lime, egg and meringue cheesecake) with no known previous reaction to vaccines. Concomitant medications included progestogen orally and an unspecified medication topically, both for menopause and Vitamin B12 orally for vitamin B12 deficiency. On 08Dec2020 during post-vaccination observation the patient developed within minutes throat closing, tongue swelling, peripheral shutdown, no wheeze, chest clear. These events were reported as anaphylaxis, and required hospitalization. Vaccinated at hospital and received vaccine as she is patient facing employee. Otherwise fit and healthy, no cardiovascular, respiratory, gastrointestinal or neurological disease. No history of allergy to medicines. History of a similar allergic reaction to lemon/lime and meringue cheesecake. After

3 mouthfuls of cheesecake, onset of reaction. Required adrenaline, ambulance and treatment as inpatient. Allergy blood tests and skin tests inconclusive (unknown what tested for). Carries Epi Pen but never used. Has remained on a gluten and dairy free diet since the reaction. On the day of vaccination, her presenting anxiety was possible allergy to eggs. Within approximately 8 minutes of vaccination, she started to cough and became hypertensive (peak 175mmHg systolic), with a heart rate (HR) of 110 beats per minute (bpm) - pulse oximetry, no trace. No wheeze, no erythema of oral mucosa, no swelling. Started clawing at her neck and described feeling of itching internally. It was reported the patient carried an adrenalin autoinjector (EPIPEN). The patient was treated with IM adrenaline, IM chlorphenamine maleate (PIRITON), IM hydrocortisone given with minimal improvement and given two nebulised adrenaline (adrenaline nebulizers) which resulted in rapid resolution of symptoms (15 minutes elapsed between administration of IM adrenaline and nebulised adrenaline). Around 20 minutes later her symptoms returned. Given nebulised adrenaline with rapid recovery. Admitted to short stay unit (emergency department (ED)) for observation and discharged around 19:30 on 08Dec2020. No tryptase testing performed, no other blood tests. There was no reaction at the injection site. On 09Dec2020 the patient was at home and reported feeling tired, with heavy limbs. She is apyrexial with no ongoing signs of allergy. Lab tests on 08Dec2020 includes: Blood pressure systolic: 175mmHg, Heart rate: 110bpm and Pulse oximetry: No trace. The patient had recovered from the events in Dec2020. The events were considered serious medically significant, for hospitalization and for being life threatening. The patient has not had symptoms associated with COVID-19. Patient has not been tested or has had an inconclusive test for COVID-19 (as reported). Patient is not enrolled in clinical trial. The vaccine was given by hospital staff member. Follow up (10Dec2020): New information received from GB-MHRA-WEBCOVID-20201209083957 and ADR 24541453-002 includes: patient history, concomitant medications, lab tests, clinical course and the only event reported was anaphylaxis.; Sender''s Comments: The reported information is limited. Based on the close temporal relationship, the subject''s signs and symptoms, being, at least in part, suggestive of anaphylaxis and the past medical history of allergy, there is a reasonable possibility that the events are related to BNT162 vaccine.

VAERS ID: 903332 (history)
Form: Version 2.0
Age: 51.0
Sex: Male
Location: North Carolina
Vaccinated: 2020-12-17
Onset: 2020-12-18
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Seizure
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Seizure approximately 24 hours after vaccination

VAERS ID: 903400 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: New York
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dyskinesia, Dyspnoea, Feeling hot, Flushing, Intensive care, Respiratory arrest, Urticaria, Wheezing
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Dyskinesia (narrow), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypersensitivity (narrow), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations: Rabies vaccine
Other Medications: Atorvastatin 20 mg once daily
Current Illness: None
Preexisting Conditions: Severe allergies
Allergies: Fish Iodine Shellfish Rabies vaccine
Diagnostic Lab Data: NA
CDC Split Type:

Write-up: 5 minutes after the Pfizer Covid-19 vaccine administration, the patient developed flushing, hives, felt warm and eventually short of breath. She started to wheeze and was wheeled into ER c/o "I can"t breathe while holding throat and thrashing with facial flushness noted. PT took 2 Benadryls and had several Epi shots. She was then discharged from the ER and later on that day, started to feel short of breath again. In the ED today she was audibly gasping for air, however had no wheezing, had a normal saturation and a normal blood pressure. She had taken another dose of her EpiPen IM and diphenhydramine 50 mg by mouth prior to coming. She was then admitted to the hospital for further observation. While on the floor, she started to feel short of breath again (about 9 am on 12/18/2020), which required an RRT . Patient received another dose of diphenhydramine IV, methylprednisolone 125 mg IV and several doses of IM epinephrine. She also required oxygen. She was then transferred to an ICU for further care.

VAERS ID: 903592 (history)
Form: Version 2.0
Age: 34.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 3/2021 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Cough, Discomfort, Dizziness, Dysphagia, Dysphonia, Dyspnoea, Hypoaesthesia, Hypoaesthesia oral, Paraesthesia, Sensation of foreign body, Tachycardia, Tachypnoea, Throat tightness, Tongue discomfort
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Asthma/bronchospasm (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad),

Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Respiratory failure (broad), Infective pneumonia (broad), Dehydration (broad), Sexual dysfunction (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations: Flu vaccine
Other Medications: Levoxyl
Current Illness: None noted
Preexisting Conditions: Graves, Hashimoto?s
Allergies: Levaquin, methimazole, flu vaccine with preservatives
Diagnostic Lab Data: Ask physicians
CDC Split Type:

Write-up: Not all or limited to: anaphylactic reaction: Feeling lump in throat, tongue feeling funny with numbness, feeling of hard to swallow, throat tightness, shortness of breath, tachycardia, tachypnea, pressure, tingling, and numbness from head to toe, dizziness/lightheartedness, cough, voice changes.

VAERS ID: 903638 (history)
Form: Version 2.0
Age: 62.0
Sex: Male
Location: Massachusetts
Vaccinated: 2020-12-17
Onset: 2020-12-18
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Feeling abnormal, Hypoaesthesia, Transient ischaemic attack, Vision blurred
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Ischaemic central nervous system vascular conditions (narrow), Dementia (broad), Embolic and thrombotic events, arterial (narrow), Guillain-Barre syndrome (broad), Glaucoma (broad), Lens disorders (broad), Retinal disorders (broad), Hypoglycaemia (broad), Sexual dysfunction (broad)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Truvada 200 mg, Protonix 40 mg, Tylenol 650 mg, ASA 81 mg, Lipitor 10 mg, Omnipaque 350mg, Lisinopril 10 mg, Zofran 4 mg
Current Illness: none
Preexisting Conditions: HTN
Allergies: NKDA
Diagnostic Lab Data: admitted to hospital TIA
CDC Split Type:

Write-up: Patient is a very pleasant 62 year old gentleman with a history of HTN, hyperlipidemia who presented with left facial numbness and left UE numbness. He states that it is worse medially on his arm. Present in upper and lower face. He states it started around 730 am this morning. He also notes intermittent "foggy" sensation since Monday associated with some blurred vision that comes and goes. Denies focal weakness, unsteady gait, difficulty with speech and swallow. He denies f/c, cp, sob, rash, pruritis, n/v/d, edema. He did receive the Pfizer COVID vaccine yesterday

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VAERS ID: 903766 (history)
Form: Version 2.0
Age: 36.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angioedema, Full blood count, Human chorionic gonadotropin, Metabolic function test
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Oropharyngeal allergic conditions (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Levothyroxine 100 mcg, albuterol HFA, Wellbutrin XL
Current Illness: None
Preexisting Conditions: CSF leak, migraines, asthma
Allergies: Amoxicillin (hives), divalproex (swelling)
Diagnostic Lab Data: CBC, BMP, HCG
CDC Split Type:

Write-up: Angioedema

VAERS ID: 903800 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Puerto Rico
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	LA / SYR

Administered by: Public **Purchased by:** ?
Symptoms: Chest X-ray, Chest pain, Dyspnoea, Electrocardiogram, Full blood count, Malaise, Metabolic function test, Nausea, Throat tightness, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No

Previous Vaccinations:
Other Medications:
Current Illness: asthma
Preexisting Conditions: asthma
Allergies: seafood, barbiturates, caffeine, oxycodone, demerol, morphine, acetaminophen, nalbuphine
Diagnostic Lab Data: cbc, bmp, ekg, Chest xray
CDC Split Type:

Write-up: sob, nausea, malaise, vomit, chest pain, throat tightness

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VAERS ID: 903806 (history)
Form: Version 2.0
Age: 89.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-17
Onset: 2020-12-18
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Lorazepam, Depakote, Trazadone, Coreg, Bumetanide, Senna, Bumetanide
Current Illness: Mild cough
Preexisting Conditions: A fib, Gastro reflux, type 2 diabetes, hypertension, edema, anxiety, heart failure, pacemaker, obesity, OA, anemia
Allergies: Codeine, morphine, propafenone
Diagnostic Lab Data: Tests performed at hospital

CDC Split Type:

Write-up: Resident became short of breath 7 hours after vaccine, went to hospital, was COVID+

VAERS ID: 903820 (history)
Form: Version 2.0
Age: 85.0
Sex: Male
Location: Florida
Vaccinated: 2020-12-17
Onset: 2020-12-18
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, Laboratory test, Pyrexia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: eye drops, hydroxyzine, Ivermectin, Xarelto, donepezil, finasteride, melatonin, Tamsulosin, atorvastatin, fluxetine
Current Illness: None
Preexisting Conditions: Dementia, anxiety, glaucoma, hyperlipidemia, COPD
Allergies: Morphine
Diagnostic Lab Data: tests performed at hospital
CDC Split Type:

Write-up: Day following vaccine, fever, sob - went to hospital, COVID+

VAERS ID: 903931 (history)
Form: Version 2.0
Age: 29.0
Sex: Male
Location: Washington
Vaccinated: 2020-12-17
Onset: 2020-12-18
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	AR / SYR

Administered by: Military **Purchased by:** ?
Symptoms: Abdominal pain, Blood creatinine increased, C-reactive protein increased, Computerised tomogram abdomen, Computerised tomogram abnormal, Dehydration, Gastrointestinal inflammation, Gastrointestinal wall thickening, Haematocrit increased, Scan with contrast abnormal, Small intestinal obstruction
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Retroperitoneal fibrosis (broad), Malignancy related therapeutic and diagnostic procedures (narrow), Gastrointestinal obstruction (narrow), Gastrointestinal nonspecific inflammation (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Ischaemic colitis (broad), Chronic kidney disease (broad), Noninfectious diarrhoea (broad), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Denies any home medications
Current Illness: None
Preexisting Conditions: None
Allergies: Denies any allergies
Diagnostic Lab Data: 12/18/20 - CT Abdomen and Pelvis w/ Contrast
CDC Split Type:

Write-up: Presented with periumbilical pain to emergency department (patient works at Medical Center). Admitted to hospital for small bowel obstruction. Labs were consistent with dehydration (Hct 55, Cr 1.25), as well as CRP 1.10. A CT Abd/Pelvis identified proximal dilation and fecalization of small bowel, with a transition point in the left lower quadrant. Distal to the transition point, the small bowel appears thick, with hyperenhancement and

inflammation progressing into the cecum. General Surgery was consulted and patient admitted to hospital for management of this small bowel obstruction.

VAERS ID: 903938 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Injection site induration, Injection site pain, Injection site rash, Injection site warmth, Rash
SMQs:, Anaphylactic reaction (broad), Extravasation events (injections, infusions and implants) (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Microgestin
Current Illness: none
Preexisting Conditions: none
Allergies: NKA
Diagnostic Lab Data: 6:11PM: 185/71 Temp 98.2 sat 98% HR85
CDC Split Type:

Write-up: rash appeared at site of injection, very painful, hot to touch, hard, with rash spreading to torso

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VAERS ID: 904072 (history)
Form: Version 2.0

Age: 34.0
Sex: Female
Location: New York
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-19

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest discomfort, Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: allegra Klonopin Lamictal Latuda Implanon PRILOSEC Synthroid albuterol inhaler
Current Illness:
Preexisting Conditions: Asthma gerd hypothyroid gastric sleeve GI bleed
Allergies: Latex, eggs, pork, shellfish, beef, cinamon
Diagnostic Lab Data:
CDC Split Type:

Write-up: Chest Tightness and shortness of breath

VAERS ID: 904107 (history)
Form: Version 2.0
Age: 82.0
Sex: Male
Location: Florida
Vaccinated: 2020-12-19
Onset: 2020-12-19
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-19

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Probiotic Capsule, Macrobid Capsule 100 MG, Simethicone Tablet 80 MG, Meloxicam Tablet 7.5 MG, Asmanex HFA Aerosol 100 MCG/ACT (Mometasone Furoate), Incruse Ellipta Aerosol Powder Breath Activated 62.5 MCG/INH (Umeclidinium Bromide), Mirta
Current Illness: SEPSIS, UNSPECIFIED ORGANISM, THROMBOCYTOPENIA, ADULT FAILURE TO THRIVE, CHRONIC OBSTRUCTIVE PULMONARY DISEASE
Preexisting Conditions: COPD, Adult failure to thrive
Allergies: NKA
Diagnostic Lab Data:
CDC Split Type:

Write-up: Resident short of breath approximately 10 minutes post vaccination. Staff noted that his baseline upon activity and transferring is generally shortness of breath. Resident wears 2L NC, O2 saturation was 92%, increased to NC to 4 Liters O2 saturation up to 96%. 170/80, heart rate 94, regular rhythm. Sent to hospital.

VAERS ID: 904234 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-19

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) /	- / 1	RA / IM

PFIZER/BIONTECH		

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Palpitations, Rash, Tachycardia
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Imitrex, Topamax
Current Illness:
Preexisting Conditions: migraines
Allergies: None
Diagnostic Lab Data:
CDC Split Type:

Write-up: Rach, tachycardia, palpitations, shortness of breath shortly after receiving vaccine - given epi, solumedrol, Benadryl. persistent symptoms the following day.

VAERS ID: 904264 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: California
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-19

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain lower, Blood triglycerides normal, Computerised tomogram abdomen, Computerised tomogram abnormal, Constipation, Lipase increased, Liver function test normal, Nausea, Pancreatitis acute, Scan with contrast abnormal, Ultrasound abdomen normal, Vomiting, White blood cell count increased

SMQs:, Acute pancreatitis (narrow), Neuroleptic malignant syndrome (broad), Retroperitoneal fibrosis (broad), Malignancy related therapeutic and diagnostic procedures (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Propranolol 60mg ER daily Sertraline 100mg daily
Current Illness: No acute illnesses at time of vaccination, unknown up to one month prior
Preexisting Conditions: Hypertension Depression
Allergies: No known medication allergies, unknown other allergies
Diagnostic Lab Data: Lipase $g 6000 IU/L on 12/18/20, 967 IU/L on 12/19/20 White Blood Cell count 20,000 on 12/18/20, 11,550 on 12/19/20 liver function tests normal on 12/18/20 CT abd/pelvis with IV contrast read at 3:18am 12/19/20: "There are inflammatory changes surrounding the pancreas consistent with acute pancreatitis. No pseudocyst, abscess, or hemorrhage." US abdomen read at 23:41 12/18/20: "No gallstones, wall thickening, or pericholecystic fluid. Negative sonographic Murphy"s sign."
CDC Split Type:

Write-up: Patient presented to the emergency department at 8:45pm on 12/18/20 with lower abdominal pain, nausea, vomiting, and constipation that started approximately 2 hours prior to presentation, at approximately 6:45pm. Her labs were significant for a lipase of $g 6000 IU/L, and a CT scan of her abdomen/pelvis was done that demonstrated evidence of acute pancreatitis. Given the fact that she does not have a history of heavy alcohol use, with normal triglycerides and no evidence of gallstones on her current admission, and no recent gastroenterology procedures, there is no clear etiology of her pancreatitis; concern for post-vaccination pancreatitis. The patient is currently admitted to the hospital, on hospital day #1 of her current condition.

VAERS ID: 904349 (history)
Form: Version 2.0
Age: 36.0
Sex: Female
Location: North Dakota
Vaccinated: 2020-12-14
Onset: 2020-12-16
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-20

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Computerised tomogram, Lipase, Nausea, Pancreatitis acute, Vomiting
SMQs:, Acute pancreatitis (narrow), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: singulair, albuterol, Adderall XR, Symbicort, clonidine, B12 supplement, omeprazole, fexofenadine, fluoxetine, Mirena, levothyroxine, lorazepam, oxcarbazepine, Phenergan, Vitamin D3
Current Illness:
Preexisting Conditions: Bipolar 1, Severe persistent Asthma, Morbid Obesity
Allergies:
Diagnostic Lab Data: lipase, CT scan
CDC Split Type:

Write-up: acute, mild pancreatitis, associated with symptoms associated with Nausea and vomiting and abdominal pain. Patient''s symptoms started 1 day after her vaccination.

VAERS ID: 904385 (history)
Form: Version 2.0
Age: 83.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-16
Onset: 2020-12-16
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-20

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Comminuted fracture, Epistaxis, Facial bones fracture, Haemorrhage, Loss of consciousness, Subcutaneous haematoma, Tenderness

SMQs:, Torsade de pointes/QT prolongation (broad), Haemorrhage terms (excl laboratory terms) (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Accidents and injuries (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: amLODIPine, 7.5 mg, oral, Nightly ? cevimeline, 30 mg, oral, TID ? cholecalciferol (vitamin D3), 2,000 Units, oral, Daily ? irbesartan-hydrochlorothiazide, 1 tablet, oral, QAM ? levothyroxine, 50 mcg, oral, Daily at 0600 ? metoprolol s
Current Illness: None
Preexisting Conditions: Sjogrens Syndrome, Hypertension, Hypothyroidism
Allergies: Quinilones, Sulfa
Diagnostic Lab Data: There is a comminuted fracture of the nasal bones bilaterally with Left orbital floor blowout fracture with hemorrhagic opacification of the left maxillary sinus and slight inferior displacement with slight extraconal fat protrusion. There is associated left orbital preseptal and postseptal emphysema. Comminuted fracture involving the anterior and posterior wall of the left maxillary sinus with hemorrhagic opacification of the sinus and left malar subcutaneous hematoma
CDC Split Type:

Write-up: Patient is a pleasant 83 y.o. female pediatrician with history of Sjogren''s, hypothyroidism, hyperlipidemia, hypertension who had been at Hospital to get her Covid vaccine. 30 minutes after doing so she reports being in the lobby and about to walk upstairs and feeling fine. The next thing she knows she wakes up on the stairs with her nose and face bleeding surrounded by healthcare team. She denies any precipitating symptoms such as chest pain, shortness of breath, fevers dizziness, headache. She reports feeling well otherwise in the last few days. I did a thorough bony palpation exam including spine and he only point of tenderness besides on her face was the area above her right ankle. She does not have a history of syncope or collapse

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VAERS ID:	904386 (history)
Form:	Version 2.0
Age:	63.0
Sex:	Female
Location:	Illinois
Vaccinated:	2020-12-19
Onset:	2020-12-19

Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-20

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Electrocardiogram abnormal, Palpitations, Sinus tachycardia, Supraventricular tachycardia
SMQs:, Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: albuterol (VENTOLIN HFA) 108 (90 Base) MCG/ACT inhaler fluticasone NASAL (FLONASE) 50 MCG/ACT nasal spray escitalopram (LEXAPRO) 20 MG tablet
Current Illness: potassium 3.4 on day of vaccination
Preexisting Conditions: Hypertension, psoriasis
Allergies: No known allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient had vaccine at 1330 on 12/20. At around 1815 she began experiencing heart palpitations. She presented to the ED and she was found to have a heart rate in the 130s. EKG showed junctional tachycardia. She was given 6mg of adenosine and an EKG was repeated and showed sinus tachycardia. Eventually her heart rate decreased to the 70s-90s. She was noted to have a potassium of 3.4 which was repleted. She was admitted overnight for observation. In the morning her potassium was normal and she remained in sinus rhythm. She was discharged later that afternoon.

VAERS ID: 904436 (history)
Form: Version 2.0
Age: 47.0
Sex: Male
Location: New Jersey
Vaccinated: 2020-12-17
Onset: 2020-12-18
Days after vaccination: 1
Submitted: 0000-00-00

Entered: 2020-12-20

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Activated partial thromboplastin time shortened, Alanine aminotransferase normal, Aspartate aminotransferase normal, Atrial fibrillation, Blood albumin normal, Blood alkaline phosphatase normal, Blood creatinine normal, Blood culture negative, Blood lactic acid decreased, Blood potassium decreased, Chest X-ray abnormal, Gram stain positive, Haematocrit decreased, Hypotension, International normalised ratio increased, Malaise, Palpitations, Platelet count normal, Procalcitonin increased, Protein total normal, Pyrexia, Respiratory tract congestion, Tachycardia, White blood cell count increased
SMQs:, Liver-related coagulation and bleeding disturbances (narrow), Anaphylactic reaction (broad), Haematopoietic erythropenia (broad), Haemorrhage laboratory terms (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (narrow), Sepsis (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Flomax 0.4mg Daily
Current Illness: Enterococcus faecalis UTI diagnosed 11/2520 associated with ureteral stent removal 11/10/20 treated successfully with Augmentin
Preexisting Conditions: Borderline HTN, recurrent renal calculi
Allergies: None
Diagnostic Lab Data: Presentation labs: WBC 21.6 Hct 38.9 Plt 227 K 3.3 Cr 1.26 Lactate 2.6 Protein 6.5 Alb 3.5 AST/ALT/AP normal INR 1.4 PTT 48.1 Procalcitonin 4.7 CXR: Mild congestion Cultures Blod, urins (1day): NGTD
CDC Split Type:

Write-up: The patient was well prior to vaccination (12/17). The day after, he felt mildly unwell and had a low grade fever. The following day, he had a fever of 102. He received 1L of fluid at Urgent Care and had a BP ion the 80s. Shortly thereafter, he felt palpitations and developed AF. He came to the hospital where he was tachycardia to 200 bpm and hypotensive to SBP70s. He received aggressive fluid resuscitation (4L), IV metoprolol and was started on empiric Abx. Within several hours, the HR lowered, BP increased, and AF spontaneously converted to sinus. He had no dysuria. Curtures so far have not shown growth at our hospital. Urinary culture from urgent care has reportedly shows 20k gram positive cocci.

VAERS ID: 904555 (history)

Form: Version 2.0
Age: 63.0
Sex: Male
Location: New York
Vaccinated: 2020-12-15
Onset: 2020-12-17
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-20

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK 5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Blood calcium increased, Blood chloride normal, Blood creatinine normal, Blood glucose normal, Blood lactic acid decreased, Blood magnesium normal, Blood potassium normal, Blood sodium normal, Blood urea normal, Carbon dioxide normal, Chest X-ray normal, Computerised tomogram head normal, Diarrhoea, Electrocardiogram normal, Electroencephalogram normal, Fatigue, Gait disturbance, Haematocrit normal, Haemoglobin normal, Injection site pain, Magnetic resonance imaging brain normal, Memory impairment, Oropharyngeal pain, Platelet count normal, Seizure like phenomena, Syncope, Vomiting projectile, White blood cell count increased
SMQs:, Torsade de pointes/QT prolongation (broad), Acute pancreatitis (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Convulsions (narrow), Pseudomembranous colitis (broad), Parkinson-like events (broad), Gastrointestinal perforation, ulcer, haemorrhage, obstruction non-specific findings/procedures (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Depression (excl suicide and self injury) (broad), Hypotonic-hyporesponsive episode (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Bystolic 5mg once a day Prilosec 20mg once a day One a Day multivitamins
Current Illness: No other illnesses at the time of vaccination or one month prior.
Preexisting Conditions: History of hypertension and GERD.
Allergies: No allergies.
Diagnostic Lab Data: 12/17/2020: Na+ 141, K+ 4.4, Cl 101, CO 24, BUN 13, Cr 0.82,

Glucose 124, Ca 10.6, Mg 1.7 WBC 11.3, HB 18.1, Hct 52.2, Plts 173 Lactic acid 4.9 (range 0.5-2.2), repeat after hydration 1.1 CXR, CT scan of brain, MRI and E of brain, and EEG all normal
CDC Split Type:

Write-up: I had no reaction following the vaccination. The next day I had very mild soreness at the injection site. The next morning (about 36 hours after the vaccination) I woke up with fatigue and a sore throat. I had breakfast and about 10 minutes later I vomited everything (projectile vomiting, no nausea or abdominal pain). An hour later I had episode of severe watery diarrhea (just one episode). Felt very weak so I decided to sit down, stumbled to a chair, and then proceeded to have a syncopal episode with about 4 minutes of seizure like activity (witnessed, I don"t remember that part). Decided to go to the ER, where I had labs, EKG, CXR, head CT scan, MRI and EEG. I was admitted for 24 hour observation, all the tests were normal.

VAERS ID: 904719 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-16
Onset: 2020-12-18
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram cerebral abnormal, Central nervous system lesion, Demyelination, Dizziness, Gastrointestinal disorder, Headache, Injection site erythema, Injection site pruritus, Limb discomfort, Magnetic resonance imaging brain abnormal, Neurological symptom, Palpitations, Paraesthesia, Vaccination site swelling
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Optic nerve disorders (broad), Cardiomyopathy (broad), Demyelination (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Vestibular disorders (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes

Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: hydroxyzine, Xyzal, nasal spray, vitamins C, D
Current Illness: vertigo
Preexisting Conditions: asthma
Allergies: benadryl
Diagnostic Lab Data: CTA brain- found infundibulum MRI brain- found lesion and demyelination lumbar puncture- still to be done ** currently admitted in the hospital when this report was given
CDC Split Type:

Write-up: Day 1: palpitations, dizziness Day 2: headache, redness, swelling, itching on vaccination site; GI disturbance Day 3: vaccination arm had paresthesia, heaviness, almost stroke-like symptoms; the symptoms started suddenly ** treated in the ER; IV steroids- for possible allergic reaction; tpa- for the stroke-like signs; halfway through the tpa, I felt an improvement with the arm

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VAERS ID: 905191 (history)
Form: Version 2.0
Age: 86.0
Sex: Male
Location: Michigan
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: COVID-19, Dyspnoea, Hospice care, Pyrexia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No

Previous Vaccinations:
Other Medications: Aspirin chewable tab, ferrous sulfate tab, memantine tab, and tiotropium inhaler
Current Illness: Inpatient admission 12/3/2020 - 12/7/2020 for acute encephalopathy secondary to acute UTI and suspected pneumonia. Discharged from hospital to long term care on 12/7/2020
Preexisting Conditions: Bladder muscle dysfunction - overactive Alzheimer"s disease History of cerebrovascular accident COPD History of prostate cancer Iron deficiency anemia
Allergies: No known medication, product, or food allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Do no suspect that vaccine caused patient condition and resulting inpatient admission. Suspect patient had COVID-19 at time of vaccination, but had not developed symptoms yet. Here is timeline: Patient went to ED on 12-18-2020 at 22:51 with complaint for fever and shortness of breath. Patient ended up testing positive for COVID-19, 12-19-2020 00:09, but was not symptomatic at time of vaccine. As of 12/21/2020 12:04 pm, patient is still inpatient and on comfort care/hospice.

VAERS ID: 905320 (history)
Form: Version 2.0
Age: 33.0
Sex: Male
Location: New York
Vaccinated: 2020-12-16
Onset: 2020-12-18
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Appendicectomy, Appendicitis, Computerised tomogram abdomen abnormal, Laparoscopic surgery, Scan with contrast abnormal
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Malignancy related therapeutic and diagnostic procedures (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No

Previous Vaccinations:
Other Medications: Norvasc 5 mg PO Daily, Vitamin C 1 tab daily
Current Illness: None
Preexisting Conditions: Hypertension, Hyperlipidemia
Allergies: NKDA
Diagnostic Lab Data: CT Scan of Abdomen with contrast on Friday December 16th confirmed early appendicitis
CDC Split Type:

Write-up: severe abdominal pain experience 2 days post vaccination of dose 1 of 2. Diagnoses with early acute appendicitis on Friday December 18th and had a laproscopic appendectomy on Saturday December 19th.

VAERS ID: 905345 (history)
Form: Version 2.0
Age: 22.0
Sex: Male
Location: Wisconsin
Vaccinated: 2020-12-17
Onset: 2020-12-20
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Haemorrhage, Platelet count decreased, SARS-CoV-2 test negative, Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None reported by patient
Current Illness: No known past medical history
Preexisting Conditions: None
Allergies: No known allergies
Diagnostic Lab Data: COVID (-) Platelets 2000 cells/mcL

CDC Split Type:

Write-up: Patient received Pfizer COVID 19 vaccine last Thursday 12/17. Admitted today (12/21) with bleeding and low platelet count - working up for ITP, TTP. Given recency of vaccination and no known contributory allergy or medical history, physician thought potentially associated with vaccination.

VAERS ID: 905774 (history)
Form: Version 2.0
Age: 57.0
Sex: Male
Location: Indiana
Vaccinated: 2020-12-18
Onset: 2020-12-19
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Myocardial infarction
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Nurse reports that patient had no problems after receiving vaccination. Patient went home and EMS was called early the next morning and team administered vaccination was contacted physician that the associate works for stating the patient had a heart attack.

VAERS ID: 905974 (history)

Form: Version 2.0
Age: 44.0
Sex: Female
Location: California
Vaccinated: 2020-12-20
Onset: 2020-12-20
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram cerebral normal, Arteriogram carotid normal, Blood glucose normal, Computerised tomogram head normal, Drug screen positive, Hemiparesis, Intensive care, Magnetic resonance imaging brain normal, Paraesthesia, Throat tightness, Tic, Urine analysis normal, White blood cell count increased
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Dyskinesia (broad), Dystonia (broad), Drug abuse and dependence (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Nexplanon birth control implant
Current Illness: None
Preexisting Conditions: Thalassemia minor
Allergies: Egg-containing compound Latex Penicillins
Diagnostic Lab Data: 12/20/20 at 1620 Blood glucose = 73 mg/dl 12/20/20 at 1620 White Blood Cells = 10.72 thousand/mm3 12/20/20 at 1758 Urinalysis: negative for infection 12/20/20 at 1758 Urine Toxicology: positive for amphetamine-methamphetamine 12/20/20 at 1636 CT of Head without contrast: negative for bleed 12/20/20 at 2018 CT Angiography of Head and Neck: no significant blockage or stenosis 12/21/20 at 1316 MRI of Brain without contrast: normal evaluation; without any evidence of acute ischemic stroke
CDC Split Type:

Write-up: Left sided weakness of face, arm and left leg, onset 15 minutes after receiving vaccination Brought immediately to ED, subjective feeling of closing of throat. Given IM

epinephrine 0.3mg x 1. Upon evaluation in the ED by tele-neurology consult, she received 88.5mg of alteplase on 12/20/20 at 1721 She was admitted to the Intensive Care Unit on 12/20/20 at 2208 Seen by neurology on 12/21/20 at 1227. Evaluation showed weakness on the left side but is noted that it could be effort-related. Neurologist noted that patient was treated with alteplase; CT angiogram showed no significant blockage or stenosis. Noted that this is could be related to a vasovagal effect, psychogenic or an acute ischemic stroke. As of 12/21/20 at 1615, attending provider noted left-sided paresthesia, left-sided tics and possible transient ischemic attack

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VAERS ID: 906282 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: Missouri
Vaccinated: 2020-12-20
Onset: 2020-12-20
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Aphasia, Dyspnoea, Hypopnoea, Intensive care, Throat clearing, Wheezing
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Asthma/bronchospasm (broad), Dementia (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hypersensitivity (broad), Respiratory failure (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: IVIG, Mestinon, Lisinopril, Immodium, Advil, Imuran, Baclofen
Current Illness:
Preexisting Conditions: Myastheniz Gravis, Multiple Sclerosis, PCOS, HTN
Allergies: IVIG, Tecfideral, Atropine
Diagnostic Lab Data: unknown

CDC Split Type:

Write-up: Given the vaccine at 712 pm on 12/20/20. At approximately 715 pm, she began to clear her throat and then became unable to speak, followed by audible wheezes and short, shallow breaths. At 1923, Epinephrine was administered. At 1928, she was able to speak again and was transported to the ED. The patient reports after arrival to the ER, the symptoms returned. She was given PO Benadryl, followed by IV Benadryl, and then a 2nd dose of Epinephrine. She was admitted to the ICU for observation.

VAERS ID: 906285 (history)
Form: Version 2.0
Age: 64.0
Sex: Male
Location: California
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	AR / IM

Administered by: Other **Purchased by:** ?
Symptoms: Acute coronary syndrome, Altered state of consciousness
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Blood thinners x 2 yrs
Current Illness: 2-3 Household members + COVID-19
Preexisting Conditions: STATUS POST STOKE 2018 (EMBOLIC)
Allergies: Unknown
Diagnostic Lab Data:
CDC Split Type:

Write-up: At work patient had ALOC x10 minutes. Rapid response called. Transf to Hospital (12/18-12/20). D/C Dx ACUTE CORONARY SYNDROME (NON-STEMI)

VAERS ID: 906064 (history)
Form: Version 2.0
Age: 31.0
Sex: Female
Location: Massachusetts
Vaccinated: 2020-12-20
Onset: 2020-12-20
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	FH9899 / 1	LA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Cold sweat, Dyspnoea, Heart rate increased, Nausea, Pruritus, Rash
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Birth control, adderall, vitamin b complex, vitamin d, multivitamin and hair skin and nails.
Current Illness: None.
Preexisting Conditions: None.
Allergies: penicillin, amoxicillin and mold.
Diagnostic Lab Data:
CDC Split Type:

Write-up: Rapid heart beat, difficulty breathing, itching, rash, clamminess, cold sweats, nausea

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VAERS ID: 906132 (history)

Form: Version 2.0
Age: 50.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Chest discomfort, Feeling hot, Tachycardia
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020501255

Write-up: waves of heat feeling; tachycardia; chest tightness; This is a spontaneous report from contactable physician via Pfizer Sales Representative. A 50-year-old female patient received bnt162b2, via an unspecified route of administration in 2020 at single dose for immunization. The patient''s medical history and concomitant medications were not reported. The patient experienced waves of heat feeling, tachycardia, chest tightness 15 min after receiving the vaccine in 2020. Patient received several doses of steroid and was kept overnight in hospital and recovered. The outcome of events was recovered in 2020. Information on the Lot/Batch number has been requested.; Sender''s Comments: The reported information is limited. Based on the close temporal relationship and the description of the events, waves of heat feeling, tachycardia, chest tightness, there is a reasonable possibility that the events are related to BNT162 vaccine.The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this reviewas well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as

appropriate.

VAERS ID: 906318 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Military **Purchased by:** ?
Symptoms: Acute myocardial infarction, Blood magnesium normal, Blood phosphorus normal, Brain natriuretic peptide increased, Chest X-ray normal, Chest discomfort, Dyspnoea, Electrocardiogram ST segment depression, Emotional distress, Fibrin D dimer normal, Full blood count normal, Headache, Hypertension, International normalised ratio increased, Metabolic function test normal, Pain, Prothrombin time prolonged, SARS-CoV-2 test negative, Troponin I increased
SMQs:, Cardiac failure (broad), Liver-related coagulation and bleeding disturbances (narrow), Anaphylactic reaction (broad), Haemorrhage laboratory terms (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Hypertension (narrow), Cardiomyopathy (broad), Depression (excl suicide and self injury) (broad), Other ischaemic heart disease (broad), Hypokalaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Advair Diskus Inhalation. Albuterol Sulfate HFA Inhalation. Amlodipine Besy-Benazepril HCl Oral. Aspirin Oral. Esomep. Esomeprazole Magnesium Oral. Ibuprofen Oral. NIFEdipine ER Oral. Nitro-Dur Transdermal. PrednISONE Oral.
Current Illness:
Preexisting Conditions: Congestive Heart Failure [Active]. Heart Disease [Active]. COPD - Chronic Obstructive Pulmonary Disease [Active]. Abnormal EKG [Active]. Hypokalemia [Active]. Lung Disease [Active]. Chest Pain [Active].
Allergies: Levaquin
Diagnostic Lab Data: EKG: EKG time: (21:33 Dec 21 2020). Normal sinus rhythm. Normal P

waves. Normal PRI. Normal QRS complex. Normal axis. Normal QT and QTc. Mild ST depression in lead II and III. The study has been interpreted contemporaneously by me. The study has been independently viewed by me. The EKG appears to be a good tracing. Interpretation time: 21:35 Dec 21 2020. EKG #2: EKG time: (00:04 Dec 22 2020). No acute process. No acute ischemia. Normal EKG. Normal sinus rhythm. Normal P waves. Normal PRI. Normal QRS complex. Normal axis. Normal ST and T waves, QT and QTc. ST-depression no longer present. The study has been interpreted contemporaneously by me. The study has been independently viewed by me. The EKG appears to be a good tracing. Interpretation time: 00:05 Dec 22 2020. Chest X-ray: No acute disease. Normal lung markings present. Normal heart size. Mediastinum normal. Great vessels normal. Soft tissues normal. No infiltrate. No fracture. No bony lesion present. Views: PA. Technique: good. The X-rays were independently viewed by me and interpreted contemporaneously by me. Laboratory Tests: (COVID: neg). CBC: CBC is normal. Chemistries: Comprehensive Metabolic Panel (Chem 14)- normal. Normal PO4- and Mg. Cardiac Labs: Troponin-I 0.18 -- $g 0.15. BNP 113. Coagulation Studies: D-dimer negative. PT 20.6. INR 1.7.
CDC Split Type:

Write-up: 57-year-old female history of hypertension, hyperlipidemia, type 2 diabetes, COPD, subsegmental PE is not on anticoagulation, multiple cardiac stents presenting with greater than 12 hrs of worsening left-sided chest pressure, headache and shortness of breath. Patient takes a daily aspirin and had no improvement of symptoms with her at-home nitroglycerin. Here afebrile, HTN, remaining vitals wnl. Non-toxic, in moderate distress 2/2 to pain. EKG with minimal ST depressions in leads II and III. Will plan for CXR and labs. Pt given zofran and morphine for pain control. Will give additional aspirin for total 324 mg in last 24 hrs. On re-evaluation, pt with mild improvement in pain. Troponin elevated at 0.18, remaining labs wnl. At this time concerned for NSTEMI, pt treated with 1 mg/kg of lovenox and MOD consulted for admission. MOD evaluated pt and cardiology was consulted. Given concerning PMHx and current hx of chest pain with findings consistent with NSTEMI, cardiology at recommended likely transfer for cardiac cath. Will pend repeat troponin and EKG for dispo decision.

VAERS ID: 906333 (history)
Form: Version 2.0
Age: 66.0
Sex: Male
Location: New York
Vaccinated: 2020-12-18
Onset: 2020-12-21
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anosmia, Condition aggravated, Cough, Decreased appetite, Fatigue, Intensive care, Pyrexia, Respiratory symptom, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Taste and smell disorders (narrow), Anticholinergic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic

infections (broad), Immune-mediated/autoimmune disorders (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: GERD
Allergies: No known Allergies
Diagnostic Lab Data: Cepheid Xpert Xpress positive for SARS-CoV-2 12/21 at 10:30AM
CDC Split Type:

Write-up: Approximately 65 hours post-vaccination patient felt profound fatigue, no appetite, and had increase in baseline chronic cough, and anosmia. Patient was admitted to the hospital on 12/21 due to worsening respiratory symptoms that required supplemental oxygen-initially 2L via nasal cannula. Patient was upgraded to ICU-level of care at 6:30PM 12/21 to receive high-flow nasal cannula, and has had one episode of fever (100.6) 12/22 at 7:00 AM.

VAERS ID: 906506 (history)
Form: Version 2.0
Age: 38.0
Sex: Female
Location: Tennessee
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Flushing, Stridor, Throat irritation
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Asthma
Allergies: Denied allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Burning in throat, flushed face and neck. Developed stridor. Benadryl 25 mg PO given, Epinephrine 0.5mg given IM, albuterol inhaler 4 puffs. Transferred to the Emergency department and admitted to the hospital.

VAERS ID: 906514 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Idaho
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / UNK	RA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Cough, Dysphonia, Intensive care, Throat irritation, Throat tightness
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none known
Current Illness: T2DM, asthma, obesity, Factor V Leiden deficiency, psoriasis

Preexisting Conditions: same as above
Allergies: Anaphylactic reaction to Zolare and Humira
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient was administered the Covid19 vaccine. She was advised to wait 30 minutes post vaccination. While the patient was waiting, she reported having an itchy throat, throat tightness, then a hoarse voice and a cough developed. This happened at about 20 minutes after she received the vaccine. The patient was assessed by the nursing and provider staff. She received an adult epi pen injection and EMS was called. Patient was taken to the ER by EMS. She reports she received two more epi injections, benadryl, and Solu Medrol. She was stabilized. Patient was discharged from the ED after several hours. She then reports a second episode of throat tightening and worsening cough at 12:30 am and was taken by ambulance to the ICU and admitted. She is still in the hospital at this time 12/22/2020.

VAERS ID: 906529 (history)
Form: Version 2.0
Age: 34.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-18
Onset: 2020-12-19
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Exposure during pregnancy, Premature delivery, Premature labour, Premature rupture of membranes
SMQs:, Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Claritin, Prenatal Vitamins, Flonase
Current Illness: no
Preexisting Conditions: no
Allergies: no

Diagnostic Lab Data: No
CDC Split Type: vsafe

Write-up: When I got the vaccination I was 32weeks pregnant and on Saturday I had spontaeous rupture of the amnotic fluids and went immediately to the hospital and was immediately given steroid, magnesium for the baby. And on Sunday around 3:45PM I got a second round of the steroids and was transferred for observation. On Monday, at 8:06am I went into early labor I delivered my baby at 33weeks gestation and she weighed 3lb 11oz. Expected Date of Delivery-2/8/2021. I was a high risk patient d/t Fibroids but have experienced no issues the entire pregnancy and my last ultrasound was 12/17 and baby was healthy with no complications at that time.

VAERS ID: 906532 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-19
Onset: 2020-12-20
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Condition aggravated, Intensive care, Seizure
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: folic acid cranberry Pepcid florastor keppra oxycarbazine
Current Illness:
Preexisting Conditions: LERS-DANLOS SYNDROME gerd tachycardia seizures cognitive communication disorder
Allergies: NKDA
Diagnostic Lab Data: resident currently at hospital admitted on 12/20/2020

CDC Split Type:

Write-up: seizure, resident sent to er and in CCU

VAERS ID: 906708 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Bronchitis, Chest discomfort, Dyspnoea, Feeling abnormal, Headache, Hypoaesthesia, Injection site paraesthesia, Pain in extremity, Pneumonia, Rash, Skin discolouration
SMQs:, Anaphylactic reaction (narrow), Peripheral neuropathy (broad), Dementia (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020502007

Write-up: bronchitis or pneumonia; pain down arm into fingers; injection site tingling; Chest felt heavy and difficulty breathing; Chest felt heavy and difficulty breathing; bronchitis or pneumonia; rash on both arms with bright purple skin; rash on both arms with bright purple skin; felt like everything was numb and like she was on drugs; felt like everything was numb

and like she was on drugs; Severe headache; This is a spontaneous report from a contactable consumer (nurse) via a Pfizer sales representative. A female patient of an unspecified age received the first dose of the bnt162b2 (BNT162B2; also reported as COVID-19 VACCINE), via an unspecified route of administration on an unspecified date at a single dose for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. On an unspecified date, the patient experienced: bronchitis or pneumonia (medically significant), pain down arm into fingers, injection site tingling, chest felt heavy and difficulty breathing, rash on both arms with bright purple skin, felt like everything was numb and like she was on drugs, and severe headache; all of which required hospitalization. The clinical course was reported as follows: The female patient (nurse) took the first dose of the vaccine and experienced pain down arm into fingers, injection site tingling. The patient"s chest felt heavy and difficulty breathing; however, the "tongue never swelled." Rapid response was called, and the patient was taken to the hospital. The patient was given a "cocktail" to treat the reaction. The patient "felt like everything was numb and like she was on drugs" The patient also felt like she had bronchitis or pneumonia. The patient also experienced a "rash on both arms with bright purple skin"; along with a severe headache. Therapeutic measures were taken as a result of pain down arm into fingers, injection site tingling, and chest felt heavy and difficulty breathing. The clinical outcome of the events was unknown. No follow-up attempt possible; information about batch/lot number cannot be obtained.

VAERS ID: 906733 (history)
Form: Version 2.0
Age: 50.0
Sex: Female
Location: Maryland
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Anaphylactic reaction
SMQs:, Anaphylactic reaction (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: yes
Diagnostic Lab Data:
CDC Split Type:

Write-up: anaphylaxis

VAERS ID: 906910 (history)
Form: Version 2.0
Age: 56.0
Sex: Male
Location: Florida
Vaccinated: 2020-12-18
Onset: 2020-12-21
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood bilirubin increased, Blood osmolarity increased, Blood urea increased, Contusion, Haemorrhage, Immune thrombocytopenia, Monocyte percentage increased, Platelet count decreased, Platelet transfusion, Thrombocytopenia
SMQs:, Acute renal failure (broad), Liver related investigations, signs and symptoms (narrow), Acute pancreatitis (broad), Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Systemic lupus erythematosus (narrow), Retroperitoneal fibrosis (broad), Biliary system related investigations, signs and symptoms (narrow), Accidents and injuries (narrow), Chronic kidney disease (broad), Hypersensitivity (narrow), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: No home medications at the time of vaccination or hospital admission
Current Illness: No pertinent past medical history prior to vaccination
Preexisting Conditions: No pertinent past medical history prior to vaccination

Allergies: No known allergies of any type
Diagnostic Lab Data: BP: 159/106 (12/22/20), Platelets 0 (12/22/20), monocytes 12.1% (12/22/20), BUN 22.3 (12/22/20), BILI 1.50 (12/22/20), calculated osmolality 297 (12/22/20). All other labs are w
CDC Split Type:

Write-up: HPI: 56 y.o. male with no pmhx c/o generalized bruising for 2 days, noticed small blood tinged spots generalized. Gradual onset, severe on severity, no alleviating or aggravating factors. Patient denies fevers, chills, N/V/D, abdominal pain. In ER: Platelet <1. Platelet transfusion in ER. Admitted for Thrombocytopenia/ITP

VAERS ID: 907022 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Angioedema, Endotracheal intubation
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal allergic conditions (narrow), Hypersensitivity (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Calcium 600 mg with D daily; Women"s Multiple vitamin daily; Vitamin D3 5,000 units daily; Zyrtec 10 mg at bedtime PRN; Celebrex 200 mg daily; Gabapentin 300 mg (3 caps daily); Melatonin 9 mg daily PRN; Zanaflex 4 mg Daily
Current Illness: Not known
Preexisting Conditions: Anxiety; Depression; Chronic Back Pain
Allergies: Morphine; Tramadol; NSAIDs; Codeine; Oxycodone; Percocet
Diagnostic Lab Data: None at this time
CDC Split Type:

Write-up: Anaphylaxis/Angioedema Patient was given EpiPen 0.3 mg IM; Methylprednisolone 125 mg once; Diphenhydramine 25 mg IV push once; Famotidine 20 mg IV push once; Dexamethasone 10 mg IV push once Patient was intubated and put on propofol and midazolam drips for sedation

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VAERS ID: 907074 (history)
Form: Version 2.0
Age: 82.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Atrial fibrillation, Condition aggravated
SMQs:, Supraventricular tachyarrhythmias (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: atrial fibrillation hx
Preexisting Conditions: atrial fibrillation hx
Allergies: NKDA
Diagnostic Lab Data:
CDC Split Type:

Write-up: EMS called after patient displayed a heart rate of 160, a little over an hour after vaccine administration. Patient was taken to the hospital and diagnosed with an episode of RVR Afib; she was admitted to the hospital.

VAERS ID: 907101 (history)
Form: Version 2.0

Age: 41.0
Sex: Female
Location: Maine
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	RA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Chest X-ray, Cough, Hyperhidrosis, Nausea, Palpitations, Speech disorder
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: SIngulair, advair, Duonebs, dupixent, albuterol, zyrtec, chlorpheniramine; wellbutrin, Sertraline;
Current Illness: had been feeling slightly under the weather two weeks ago; no current asthma exacerbation, but has very labile asthma
Preexisting Conditions: Sleep apnea on CPAP eosinophilic asthma on multiple controller meds environmental allergies obesity
Allergies: Anaphylaxis hx to NSAIDS; hs of environmental allergies, and reaction to latex
Diagnostic Lab Data: 12/22/20 - CXR ordered - results not currently available
CDC Split Type:

Write-up: patient felt slightly nauseated at 10 minutes after injection, then developed slight sweating; BP 160/81; 83 at 5:45 and then 158/87 with HR 82 at 5: 52 pm. Her lungs were clear, she was speaking in full sentences and was denying any chest pressure, her usual sense of asthma exacerbation. At 6:05 it was 164/83 with HR 79 and patient developed a dry cough; we decided to have her wait just a bit longer, then cough worsened, so at 6:25, decision was made to have patient seen in ER for further assessment, and en route in wheelchair to ER the dry cough became persistent, spasmodic and patient was unable to speak. Epi-Pen was injected in right mid thigh, and patient transported to ED urgent eval. She noted immediate palpitations, and slight improvement of breahting, was able to speak in four word sentences. On arrival to the ED, patient was administered Duonebs, Albuterol neb, IV

Benedryl, IV Solumedrol; CXR was obtained, with results pending. Patient was sent to observation for ongoing monitoring and assessment of breathing. at 6:30 PM in the ER, she

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VAERS ID: 907330 (history)
Form: Version 2.0
Age: 50.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / SYR
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal discomfort, Abdominal pain, Dizziness, Dyspnoea, Headache, Hyperhidrosis, Hyperventilation, Rash macular, Tremor, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Parkinson-like events (broad), Gastrointestinal perforation, ulcer, haemorrhage, obstruction non-specific findings/procedures (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Flexeril, Advil, Protonix, Estrace, Vivelle DOT Norco PRN
Current Illness: No
Preexisting Conditions: Mild Osteopenia, GERD, Anemia,
Allergies: Reglan - anaphylaxis
Diagnostic Lab Data: 12/22 ED administrated PEPCID, ZOFRAN, Prednisone 60 mg, and Ativan 1mg IV.
CDC Split Type:

Write-up: Employee presented to COVID Clinic for Moderna COVID 19 vaccination 1st dose. Given to left arm. Left clinic prior to completing 15 minute observation time and told an MA in waiting area that she felt ill ot her stomach and having trouble taking deep breaths. Employee found in nearby Bathroom sitting on the floor, she had vomited, reported she was lightheaded, couldn"t breath, shaking, abdominal discomfort sweating, attempted to move employee to wheelchair, did respond well to transfer to Wheelchair, She reported symptoms worsening: HA, abdominal pain and developed blotchy skin, hyperventilating, and dizzy. CODE Blue called, patient given Epinephrine injection 0.5mg patient sent to ER

VAERS ID: 907423 (history)
Form: Version 2.0
Age: 26.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Chest X-ray, Computerised tomogram head, Dyspnoea, Electrocardiogram, Hypoaesthesia, Laboratory test, Nausea, Pregnancy test, Speech disorder, Swollen tongue
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (narrow), Peripheral neuropathy (broad), Dementia (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Psychosis and psychotic disorders (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: Laparoscopic Cholecystectomy on 11-6-2020
Preexisting Conditions: None
Allergies: Keflex and Tramadol

Diagnostic Lab Data: CT of head, routine labs, chest x-ray, pregnancy test, neurology consulting, EKG
CDC Split Type:

Write-up: Onset of symptoms began approximately 1.5 hours after vaccination. Patient became weak c/o tongue swelling, nausea, difficulty breathing, numbness especially in lower extremities. Felt she couldn''t breath, numbness continued to get worse and affected her upper extremities as well, weak speech.

VAERS ID: 907426 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: New Mexico
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	- / IM

Administered by: Other **Purchased by:** ?
Symptoms: Acute kidney injury, Anxiety, Atrial tachycardia, Chest discomfort, Chest pain, Dizziness, Dyspnoea, Flushing, Hyperhidrosis, Hypersensitivity, Hypertension, Malaise, Palpitations, Tremor
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Anaphylactic reaction (narrow), Angioedema (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Parkinson-like events (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hypertension (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: ? Atrial paroxysmal tachycardia ? History of repair of atrial septal defect
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: 48 y.o female with history of atrial tachycardia who presents to the ED via EMS for a possible adverse reaction to Covid vaccine. Patient received her Covid vaccine around 1600 today, and soon became diaphoretic, shaky, and lightheaded. She had presented to the emergency department. Currently she denies any chest pain, difficulty breathing, throat swelling, tongue swelling, or any other symptoms currently except for palpitations. ?Allergic Reaction The primary symptoms are shortness of breath. The primary symptoms do not include cough, abdominal pain, vomiting, dizziness or rash. The current episode started 1 to 2 hours ago. The problem has not changed since onset. The onset of the reaction was associated with a new medication. Significant symptoms also include flushing. 48-year-old female with a history of atrial septal defect, status post atrial septal defect repair in 1980. She works as a nurse at Hospital and has been experiencing increasing rapid palpitations associated with chest pain, and hypertension. With her episodes, she experiences marked lightheadedness, dyspnea, and feeling marked anxiety, as well as chest tightness. She received the COVID-19 vaccine today and while waiting in the observation room,she started feeling unwell, with rapid palpitations, associated with lightheadedness and dyspnea. She last had a sustained episode 2 - 3 weeks ago and had presented to ER and Hospital. No syncope. No orthopnea, PND or increased lower extremity swelling. Active Hospital Problems ? Diagnosis ? Atrial paroxysmal tachycardia ? History of repair of atrial septal defect ? ? ? 1. Paroxysmal atrial tachycardia in setting of prior atrial septal defect repair - she is having breakthrough episodes through flecainide/digoxin - it is likely her atrial tachycardia is related to her ASD patch. Will hold flecainide/digoxin for now, and try to schedule an ablation during her hospital admission due to highly symptomatic episodes resulting in multiple ER visits. ? 2. Acute renal insufficiency - most likely pre-renal - iv fluids started. ? 3. Possible COVID-19 vaccine reaction - she probably had an incidental atrial tachycardia episode post vaccine administration, rather than an actual adverse reaction. Continue to monitor.

VAERS ID: 907478 (history)
Form: Version 2.0
Age: 67.0
Sex: Male
Location: Indiana
Vaccinated: 2020-12-18
Onset: 2020-12-20
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram head normal, Computerised tomogram thorax normal, Electroencephalogram, Generalised tonic-clonic seizure
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Atorvastatin, Multivitamin, Aspirin
Current Illness: none
Preexisting Conditions: none
Allergies: none
Diagnostic Lab Data: CT Head, MRI Brain, CT Chest, all negative, EEG results unknown at this time
CDC Split Type:

Write-up: Seizure (Grand mal)

VAERS ID: 907616 (history)
Form: Version 2.0
Age: 53.0
Sex: Female
Location: New Mexico
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	- / IM

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Dyspnoea, Hypersensitivity, Pruritus, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Past Medical History: Diagnosis Date ? Addison''s disease (CMS-HCC) 03/2015 ? Asthma (CMS-HCC) ? Disorder of thyroid (CMS-HCC)
Allergies: meperidine, morphine=anaphylaxis flagyl=pancreatitis
Diagnostic Lab Data:
CDC Split Type:

Write-up: 12-22 HPI 53-year-old female with a history of Addison''s disease, anaphylactic reaction who presents to the ED complaining of hives and shortness of breath. Patient reports that 3 days ago she received the COVID-19 pfizer vaccine. She reports that since that time she has developed progressively worsening hives on her legs and arms. Approximately 1 hour ago she began to develop shortness of breath and so she presented to the ER. Patient reports a previous history of anaphylactic reactions multiple times. Denies any other acute complaints at this time. MDM Patient came in with shortness of breath and hives. Suspect allergic reaction to the COVID-19 vaccine. Patient had already taken 50 mg of Benadryl. She was given Solu-Medrol and EpiPen. She reported feeling better with improvement in the pruritus. She reports that she has had rebound reaction requiring EpiPen at 24 hours. Given the distance that she lives from adequate medical care and the possibility for recurrent severe reactions, the patient will be hospitalized for further observation. 12-23 Female with history of asthma and addison''s had anaphylaxis to covid vaccine. Admitted over night to ensure that she did not rebound. Received IV Dex and this am has had no reoccurrence of hives or shortness of breath. Will discharge home on epipen, hydrocortisone prn, prednisone bid for 5 days. Return to ER or go to PCP for worsening symptoms.

VAERS ID: 907635 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: Delaware
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Swollen tongue, Throat irritation
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Nitrile
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient received covid 19 vaccine. She began to experience itching throat and swollen tongue. She was sent to the Emergency Department. She received IV Benadryl 50 mg, IV famotidine 20 mg and 125 mg IV solu-medrol. Around 930, patient stated that symptoms had resolved, except for tongue being slightly swollen. Patient was admitted to the observation unit of the hospital.

VAERS ID: 907805 (history)
Form: Version 2.0
Age: 52.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-15
Onset: 2020-12-16
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Atrial fibrillation, Blood parathyroid hormone increased, Chest discomfort, Chest pain, Chills, Computerised tomogram thorax normal, Dyspnoea, Echocardiogram abnormal, Fatigue, Fibrin D dimer increased, Pain, Painful respiration, Palpitations, Tricuspid valve incompetence, Troponin normal

SMQs:, Anaphylactic reaction (broad), Haemorrhage laboratory terms (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Chronic kidney disease (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ziac, protonix, loratadine, crestor, multivitamin
Current Illness: no
Preexisting Conditions: HTN, hypercholesterolemia , primary hyperparathyroidism
Allergies: no known allergies
Diagnostic Lab Data: 12/18/20 D Dimer was elevated. CT chest was negative for PE. Troponins were normal. routine labs were performed. PTH 114. 12/19/20 Echo was normal with mild tricuspid regurgitation
CDC Split Type:

Write-up: First night following after vaccine I woke up with chest pain (i though pleuritic) which went away. I had mild body aches and fatigue, chills. next day I experienced chest(again I thought pleuritic) discomfort especially when taking a deep breath. i felt better then had mild fatigue and body aches again. day 3 post vaccine I woke up with discomfort when taking a deep breath with continued discomfort. i felt tired through the day. Then that evening i developed SOB, severe palpitations and chest pain and went to ER. Diagnosis New onset rapid A fib. I was hospitalized and once my work up was finished and I had normal sinus rhythm I was discharged home the next evening.

VAERS ID: 907852 (history)
Form: Version 2.0
Age: 83.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-21
Onset: 2020-12-23
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?

Symptoms: Diarrhoea, Headache, Nausea, Vomiting
SMQs:, Acute pancreatitis (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Allopurinol 300mg, atorvastatin 20mg, Vitamin D3 1,00U, digoxin 125mcg, Colace 100mg, metoprolol XL 100mg , Pantoprazole 40mg
Current Illness: COVID 19 PNA 11/3/20
Preexisting Conditions: HTN, Osteoarthritis, HLD, Gout, Glaucoma, DM2- diet controlled, cardiomyopathy, systolic CHF, PAF
Allergies: Alendronate, Calcitonin, Penicilinn, Raloxifene, Bimatoprost, Erythromtcin
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient received Pfizer COVID 19 vaccine on 12/21/2020. Patient felt fine that day and all day of 12/22/2020. Around 2AM on 12/23/2020, Patient woke up with a headache and started to have nausea, vomiting, and diarrhea.

VAERS ID: 907860 (history)
Form: Version 2.0
Age: 66.0
Sex: Female
Location: Unknown
Vaccinated: 2020-12-22
Onset: 2020-12-23
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Chest discomfort, Dizziness, Troponin increased
SMQs:, Anaphylactic reaction (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Vestibular disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: increased troponin.
CDC Split Type:

Write-up: Chest pressure, dizziness, increased troponin lab value.

VAERS ID: 908003 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: Michigan
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	ELO140 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Chest discomfort, Dyspnoea, Electrocardiogram, Tachycardia, Throat tightness
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: Ibuprophen
Current Illness: none
Preexisting Conditions: none
Allergies: Gadolinium MRI dye
Diagnostic Lab Data: EPINEPHRINE, SOLUMEDROL, BENADRYL, PEPCID, EKG (ST). 12/21 3:14PM REPEAT EVENT 12/22 WITH ADMISSION TO OBS UNIT FOR OBSERVATION
CDC Split Type:

Write-up: ANAPHLACTIC REACTION, SOB, CHEST PRESSURE, TIGHTNESS IN THROAT, TACHYCARDIA

VAERS ID: 908262 (history)
Form: Version 2.0
Age:
Sex: Female
Location: New York
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Intensive care
SMQs:, Anaphylactic reaction (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Allergic eosinophilia; Fish allergy; Iodine allergy; Shellfish allergy
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC2020502832

Write-up: Anaphylaxis; This is a spontaneous report from a contactable pharmacist. A 55-year-old female patient received the bnt162b2 (BNT162B2; also reported as: PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 17Dec2020 at a single dose for COVID-19 immunization. The patient"s medical history included eosinophil process allergic reaction, fish, iodine and shellfish allergy; all from an unknown date and unknown if ongoing. Concomitant medications were not reported. The patient previously took rabies vaccine for immunization and experienced anaphylactic reaction on an unspecified date. On 17Dec2020, the patient experienced anaphylaxis; which required hospitalization, and was assessed as medically significant. The patient was hospitalized for anaphylaxis from 18Dec2020 to an unknown date. The clinical course was reported as follows: The pharmacist called about a patient who received the COVID-19 vaccine on 17Dec2020 and started having a reaction approximately 30 minutes later. The patient used epinephrine (EPIPEN) and 50 mg of diphenhydramine hydrochloride (BENADRYL) and returned to the hospital on 18Dec2020. The patient was currently in the intensive care unit (ICU) receiving an epinephrine drip. The patient had a previous history of an anaphylactic reaction to the rabies vaccine, eosinophil process allergic reaction, fish, iodine and shellfish allergy. The patient was stabilized but continued to have reactions (not specified). The pharmacist had not seen the patient and was reaching out to Pfizer on behalf of the physicians. The pharmacist believed this had been reported by the hospital. The pharmacist had no patient information. Therapeutic measures were taken as a result of anaphylaxis. The clinical outcome of the event, anaphylaxis, was unknown. The batch/lot numbers for the vaccine, BNT162B2, were not provided and will be requested during follow up.; Sender"s Comments: Based on the information available, a possible contributory role of the suspect products cannot be excluded for the reported event anaphylaxis due to temporal association. However patient previous history of allergic reaction cannot be excluded to have played a contributory role

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VAERS ID: 908401 (history)
Form: Version 2.0
Age: 24.0
Sex: Female
Location: Kentucky
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: C-reactive protein increased, Chills, Dizziness, Electrocardiogram abnormal, Fibrin D dimer normal, Sinus tachycardia, Tremor
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Parkinson-like events (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Junel Birth Control
Current Illness: COVID-19 Positive with past 30 days, symptoms resolved
Preexisting Conditions: anxiety, heart murmur
Allergies: No known allergies
Diagnostic Lab Data: D-Dimer 0.64 C Reactive Protein 6.98 EKG- Sinus Tachycardia 146-160"s
CDC Split Type:

Write-up: Vaccine administered at 0730. Pt went home and went to bed. Woke up at 0930 with shaking, chills and feeling like she was going to pass out. Evaluated in ER and noted to have sinus tachycardia in 140-160"s. Given 5mg IV Lopressor which took heart rate down to 110"s. Admitted to hospital 12/23/2020.

VAERS ID: 908448 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Illinois
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM
UNK: VACCINE NOT SPECIFIED (NO BRAND NAME) / UNKNOWN MANUFACTURER	- / UNK	LA / -

Administered by: Private **Purchased by:** ?
Symptoms: Blood test, Hypoaesthesia, Influenza virus test negative, Magnetic resonance imaging normal, Muscular weakness, SARS-CoV-2 test negative, Streptococcus test negative

SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), COVID-19 (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vitamin D 50000 iu
Current Illness: None
Preexisting Conditions: Christi and syringohydromyelia, l eye scotoma
Allergies: Iodinated contrast dye dicloxicillin lidocaine nectar tape smo
Diagnostic Lab Data: Step test rapid COVID regular COVID influenza all negative 12/22/20 MRI 12/23/20 Blood work 12/23/20
CDC Split Type:

Write-up: Left arm leg and face numbness bilateral legweakness Vaccine given at 915 am Symptoms started at 11 am Called Pfizer to report at 1151 am Saw primary doctor at 2 pm Return to primary md again 12/22/20 Referral to neurosurgeon 11/22/20 Hospitalized 11/22/20 to 11/24/20 lepto Meningeal inflammation hospital

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VAERS ID: 908663 (history)
Form: Version 2.0
Age: 26.0
Sex: Female
Location: California
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Erythema, Feeling hot, Lung disorder, Paraesthesia, Pharyngeal swelling
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Accutane Multivitamin
Current Illness: Acne has improved
Preexisting Conditions: Acne
Allergies: None
Diagnostic Lab Data:
CDC Split Type:

Write-up: Within minutes I had lower chest pain and wired sensation in my lungs. My throat was swelling. I was very hot and red. I the gurney my right arm was tingling. Opposite arm from vaccine.

VAERS ID: 908685 (history)
Form: Version 2.0
Age: 22.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Pharyngeal swelling, Supraventricular tachycardia
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Supraventricular tachyarrhythmias (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Nuvaring
Current Illness: Chronic Tonsillitis Dysmenorrhea
Preexisting Conditions: N/A
Allergies: N.K.A
Diagnostic Lab Data:

CDC Split Type: vsafe

Write-up: throat swelling, SVT

VAERS ID: 908869 (history)
Form: Version 2.0
Age: 73.0
Sex: Male
Location: Oklahoma
Vaccinated: 2020-12-18
Onset: 2020-12-19
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood bilirubin increased, Blood creatinine increased, Contusion, Immunoglobulin therapy, International normalised ratio normal, Metabolic function test normal, Petechiae, Platelet count decreased, Platelet transfusion, Prothrombin time normal
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Liver related investigations, signs and symptoms (narrow), Acute pancreatitis (broad), Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Retroperitoneal fibrosis (broad), Biliary system related investigations, signs and symptoms (narrow), Accidents and injuries (narrow), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Comprehensive list displayed in additional information below. ascorbic acid daily, atorvastatin 40mg daily, cholecalciferol daily, docusate 100mg daily, levothyroxine 100mcg daily, losartan 100mg daily, zinc 50mg daily
Current Illness: Vague symptoms (anorexia, fatigue) 11/29/2020 - 12/6/2020; Received high-dose influenza vaccination on 11/25/2020 at employee health department. On 11/29/2020 patient began having generalized fatigue, anorexia, and occasional chills. Presented to Urgent care on 12/4/2020 and reported same symptoms for past 5 days with 10-12 lb weight loss and worsening fatigue. Pfizer rapid PCR COVID test negative. Patient''s symptoms improved and he returned to normal activities of daily living until 12/19/2020, 1 day after COVID19

vaccination.
Preexisting Conditions: HTN, Type II Diabetes
Allergies: NKDA
Diagnostic Lab Data: 12/21/2020 CBC: platelets of 1,000
CDC Split Type:

Write-up: 12/18/2020: COVID19 vaccine received. 12/19/2020: Patient noticed petechiae/bruising on arms, legs and face. Worsened over next 48 hours. 12/21/2020: Patient had blood drawn (CMP, PT/INR, CBC) at lab. 12/22/2020: Labs resulted; CMP and PT/INR WNL (exceptions: SCr 1.24, TBil 1.7); CBC with platelet count of 1,000 resulting in patient admission to Hospital. At admission he received 80 mg of prednisone, 40 g of IV Ig and a unit of platelets. 12/23/2020: Continued hospitalization. Patient"s platelets improved to 20,000 and he received another 35g of IV Ig. 12/24/2020: Patient discharged with platelets of 38,000.

VAERS ID: 908877 (history)
Form: Version 2.0
Age: 25.0
Sex: Male
Location: New Jersey
Vaccinated: 2020-12-24
Onset: 2020-12-24
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Hypotension, Loss of consciousness, Unresponsive to stimuli
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: no
Current Illness: no
Preexisting Conditions: no
Allergies: no
Diagnostic Lab Data: yes
CDC Split Type:

Write-up: He was fine ate lunch, in a room with a patient, felt light headed and dizziness, passed out, he became unresponsive, he was hypotensive, he is now in the ER.

VAERS ID: 908893 (history)
Form: Version 2.0
Age: 77.0
Sex: Male
Location: New York
Vaccinated: 2020-12-23
Onset: 2020-12-24
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Acute kidney injury, Blood creatine increased, Blood potassium increased, Blood urea increased, Chronic kidney disease, Electrocardiogram T wave peaked, Haemodialysis, Hyperkalaemia, Mental status changes, Pyrexia
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Chronic kidney disease (narrow), Tumour lysis syndrome (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: amlodipine 10mg QD, atorvastating 10mg QD, brimonidione-timolol, cakcium acetate 667, plavix, diltiazem 30mg BID, hyralazine 100mg TID, labetalol 200mg BID, keppra 500mg , pantoprazole 20mg, latanoprost
Current Illness: Unknown
Preexisting Conditions: Hypertension, T2 Diabetes Mellitus, End Stage Renal Disease, Heart Failure with preserved ejection fraction, Coronary Artery disease s/p stent placement, COVID Pneumonia 4/2020
Allergies: NKDA
Diagnostic Lab Data: Temp 100.8, Potassium 7.8, BUN 35, Cr 8.81
CDC Split Type:

Write-up: Patient vaccinated at Nursing home. Transferred to ER the following day when patient developed fever and altered mental status. Found to have acute kidney injury on chronic kidney disease, hyperkalemia. Required emergent hemodialysis for hyperkalemia with ECG findings of peaked T waves.

VAERS ID: 908917 (history)
Form: Version 2.0
Age: 47.0
Sex: Female
Location: Nebraska
Vaccinated: 2020-12-23
Onset: 2020-12-24
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Activated partial thromboplastin time shortened, Asthenia, Binocular eye movement disorder, Blood glucose normal, Dysarthria, Facial paralysis, Feeling abnormal, Gait disturbance, Grip strength decreased, Haemoglobin normal, International normalised ratio normal, Mobility decreased, Platelet count normal, Red blood cell count normal, Visual impairment, White blood cell count normal
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Dementia (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Glaucoma (broad), Optic nerve disorders (broad), Lens disorders (broad), Retinal disorders (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hearing impairment (broad), Ocular motility disorders (narrow), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes

Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Albuterol inhaler, Vitamin D3, citalopram, fluticasone nasal spray, tramadol, nicotine transdermal patch
Current Illness: No recent illnesses in the last 30 days
Preexisting Conditions: Tobacco dependency, depression, chronic low back pain, anxiety, vitamin D deficiency
Allergies: No known drug or food allergies
Diagnostic Lab Data: WBC: 8.1 RBC: 4.81 Hemoglobin: 15.2 Platelet Count: 188 INR: 0.9 PTT: 32 Blood Glucose: 108
CDC Split Type:

Write-up: Patient is a 47 y.o. female who arrived by Car presented to the emergency department for Stroke symptoms. Patient awoke at 6:15 this morning, some difficulty seeing out of the right eye and also was stumbling towards the left and to table. Concerned about things not being right so brought to the emergency department. Patient feels her speaking and swallowing are okay. She did drink a bit of coffee earlier. She denies headache or significant vision problems presently. Continues to not feel normal on her left side. No history of stroke and parents or siblings. She does give personal history of an occipital migraine many years ago at which time she did not have a headache but had some vision troubles. Physical Exam Vitals signs and nursing note reviewed. Constitutional: General: She is not in acute distress. Appearance: She is not ill-appearing or diaphoretic. HENT: Head: Normocephalic and atraumatic. Right Ear: Tympanic membrane normal. Left Ear: Tympanic membrane normal. Nose: Nose normal. Mouth/Throat: Mouth: Mucous membranes are moist. Pharynx: No oropharyngeal exudate or posterior oropharyngeal erythema. Eyes: Conjunctiva/sclera: Conjunctivae normal. Pupils: Pupils are equal, round, and reactive to light. Comments: Patient displays absence of left lateral movement Neck: Musculoskeletal: Normal range of motion. No muscular tenderness. Cardiovascular: Rate and Rhythm: Normal rate and regular rhythm. Heart sounds: No murmur. Pulmonary: Effort: Pulmonary effort is normal. Breath sounds: Normal breath sounds. Abdominal: General: Bowel sounds are normal. There is no distension. Palpations: Abdomen is soft. Tenderness: There is no abdominal tenderness. Musculoskeletal: Right lower leg: No edema. Left lower leg: No edema. Lymphadenopathy: Cervical: No cervical adenopathy. Skin: Findings: No rash. Neurological: Mental Status: She is alert. Cranial Nerves: Cranial nerve deficit (left facial droop, dysarthria) present. Comments: Patient"s speech seems a bit slurred to me. Absence of ocular movements towards left noted as well as upward movements. Tongue is midline. Patient is unable to shrug the left shoulder or lift the left arm off the bed. Grip strength is 4 out of 5 on the left. Left leg strength is 3 out of 5. Extremity strength on right arm and leg is 5 out of 5. After consultation with a neurologist, the patient is being transferred from the ED.

VAERS ID:	908918 (history)
Form:	Version 2.0
Age:	61.0
Sex:	Male
Location:	New York
Vaccinated:	2020-12-23
Onset:	2020-12-23
Days after vaccination:	0
Submitted:	0000-00-00

Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	RA / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood glucose decreased, Bradycardia, Heart rate decreased
SMQs:, Arrhythmia related investigations, signs and symptoms (broad), Hypoglycaemia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness: Unknown
Preexisting Conditions: Bradycardia
Allergies: No known allergies
Diagnostic Lab Data: Unknown
CDC Split Type:

Write-up: This resident received the first dose of the Pfizer Covid vaccine at the Covid vaccine clinic at medical Facility. He had a rapid response, experienced low blood glucose and low heart rate. NP and medical director were in attendance. Heart rate remained low despite all measures. He was transferred to Hospital for bradycardia.

VAERS ID: 908969 (history)
Form: Version 2.0
Age: 48.0
Sex: Male
Location: Illinois
Vaccinated: 2020-12-19
Onset: 2020-12-21
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Echocardiogram, Electrocardiogram abnormal, Laboratory test normal, Nausea,

Pain, Palpitations, Pyrexia, Scan myocardial perfusion, Ventricular arrhythmia, Ventricular extrasystoles
SMQs:, Torsade de pointes/QT prolongation (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Ventricular tachyarrhythmias (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Aspirin 81mg
Current Illness: None
Preexisting Conditions: Myasthenia Gravis
Allergies: Vancomycin
Diagnostic Lab Data: Complete lab work Nuclear stress test Echo cardiogram Multiple EKG"s
CDC Split Type:

Write-up: Starting that night 12/19/20 at approx 8pm I became nauseous with a 100.8f fever. I felt achy for the 2 days with the fever subsided. On Monday 12/21/20 I was at work (Firefighter) when I started to have palpitations. I was able to obtain a 12 lead EKG right away which showed a ventricular rhythm lasting for approx 5 min. After that I was having multiple PVC"s / Ventricular beats. I was taken to Hospital where I was admitted. Over night the rhythm subsided, all my labs were good with other test"s being done. I was discharged home. I have not had any other episodes since Monday. I was cleared by our Occ Health Doc who suggested I fill this out.

VAERS ID: 908973 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-19
Onset: 2020-12-19
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) /	- / UNK	- / -

PFIZER/BIONTECH		

Administered by: Private **Purchased by:** ?
Symptoms: Burning sensation, Dizziness, Hypoaesthesia, Palpitations, Paraesthesia
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Sexual dysfunction (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: 15 min after receiving Covid 19 vaccine patient started to feel like her heart was racing / felt faint. Burning feeling in upper thigh and pelvic area. BP 180/100 HR 130. Rapid Response called / transported to ER. Admitted for 24 hr observation.. Solu -medrol, Benadryl and Ativan given in ER. Released home the next day. 72 hrs later patient states she has numbness and tingling in hands and feet. 12/24/2020 patient reports she is feeling better today / no symptoms noted.

VAERS ID: 909031 (history)
Form: Version 2.0
Age: 87.0
Sex: Female
Location: Alaska
Vaccinated: 2020-12-18
Onset: 2020-12-24
Days after vaccination: 6
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood bilirubin increased, Blood creatinine increased, Blood lactic acid decreased, C-reactive protein increased, Confusional state, Hypotension, Platelet count

decreased, SARS-CoV-2 test negative, Sepsis, White blood cell count increased
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Liver related investigations, signs and symptoms (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Haematopoietic thrombocytopenia (narrow), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Dementia (broad), Biliary system related investigations, signs and symptoms (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (broad), Sepsis (narrow), Opportunistic infections (broad), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Levothyroxine, vitamin D, Naproxen, Namenda, Aricept
Current Illness: UTI
Preexisting Conditions: Dementia
Allergies: None
Diagnostic Lab Data: Lactic acid 2.2 Creat 1.2 CRP 8.32 T. Bili 1.1 WBC 13.3 Sars Cov2 negative (rapid Abbott) PLT 115
CDC Split Type:

Write-up: Patient presented with signs and symptoms of sepsis, developing over 12 to 24 hours 6 days after vaccination. was hypotensive and confused (beyond baseline)

VAERS ID: 908243 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-11
Onset: 2020-12-11
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ 0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Feeling hot, Heart rate, Paraesthesia oral, Rash erythematous, Sinus

tachycardia
SMQs:, Anaphylactic reaction (broad), Supraventricular tachyarrhythmias (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Patient has not had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial. Patient is not pregnant.
Allergies:
Diagnostic Lab Data: Test Date: 20201211; Test Name: Pulse rate; Result Unstructured Data: Test Result:125
CDC Split Type: GBPFIZER INC2020506818

Write-up: Localised feeling of warmth; Sinus tachycardia; Red rash; Tingling lips; This is a spontaneous report from a contactable other hcp downloaded from the Medicines Agency (EMA) Regulatory Authority-WEB GB-MHRA-ADR 24542591, WWID number GB-MHRA-WEBCOVID-20201211184846. A 33-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, batch/lot number Ej 0553 , with unknown expiration date), via an unspecified route of administration on 11Dec2020 15:30 at single dose for covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. The patient experienced localised feeling of warmth, sinus tachycardia, red rash, neck rash, tingling lips on 11Dec2020. The patient underwent lab tests and procedures which included heart rate: 125 on 11Dec2020. Patient had not tested positive for COVID-19 since having the vaccine. Patient had not had symptoms associated with COVID-19. Patient had not been tested/or had had an inconclusive test for COVID-19. Patient was not enrolled in clinical trial. Patient was not pregnant. The patient received the vaccine at 15:30 within 10 minutes she had a hot sensation in her arm which developed into a red rash, this spread to her other arm. She developed a rash to both her chest and back. Her pulse was 125 and developed a strange tingling sensation to her inner bottom lip. She worked in the emergency department so knew she was having a reaction. She received chlorphenamine maleate, intravenous (IV) hydrocortisone and IV fluids. After 3 hours her symptoms settled but remained shaky. She now had steroids and antihistamines for the next three days. The outcome of localised feeling of warmth, sinus tachycardia, tingling lips was resolved. The outcome of Red rash was resolving. No follow-up attempts are possible. No further information is expected.

VAERS ID: 908247 (history)
Form: Version 2.0

Age:
Sex: Unknown
Location: Foreign
Vaccinated: 2020-12-14
Onset: 2020-12-14
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Epistaxis, Facial pain, Hypoaesthesia, Hypoaesthesia oral, Oral pain, Pain of skin, Paraesthesia, SARS-CoV-2 test, Skin warm, Tinnitus
SMQs:, Peripheral neuropathy (broad), Haemorrhage terms (excl laboratory terms) (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Glaucoma (broad), Hearing impairment (narrow), COVID-19 (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Test Result: Negative ; Comments: No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2020507315

Write-up: Numb mouth; Skin warm; Facial pain; Mouth pain; sore skin; Numbness in leg/numb tingling hands, feet, ankles, face, legs; Numbness facial; Nose bleeds; Worsening of tinnitus; Tingling feet/hands/tingling hands, feet, ankles, face, legs; This is a spontaneous report from a contactable consumer downloaded from the Medicines Agency (MA) Regulatory Authority-WEB GB-MHRA-WEBCOVID-20201214220957 Safety Report Unique Identifier GB-MHRA-ADR 24543227. EU Local Number was EU-EC-10007206182. Worldwide Unique Case Identification was GB-MHRA-WEBCOVID-20201214220957. A patient of unspecified age and gender started to receive BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on 14Dec2020 at single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. Patient had not had symptoms associated with COVID-19. Patient was not enrolled in clinical trial. The patient experienced numbness facial on 14Dec2020, nose bleeds on 14Dec2020, worsening of tinnitus on 14Dec2020, numb mouth on an unspecified date, skin warm on an

unspecified date, facial pain on an unspecified date, tingling feet/hands on an unspecified date, mouth pain on an unspecified date, and numbness in leg on 14Dec2020. The events were serious with hospitalization. As of 22Dec2020, it was also reported that patient experienced mouth and face numb sore, numb tingling hands, feet, ankles, face, legs, sore skin, and hot skin. Lab data included COVID-19 virus test was negative. Patient has not tested positive for COVID-19 since having the vaccine. Outcome of numbness facial, tingling feet/hands, and numbness in leg was not recovered. Outcome of nose bleeds was of recovered with sequelae. Outcome of worsening of tinnitus was not recovered. Outcome of numb mouth, skin warm, facial pain, mouth pain was unknown. No follow-up attempts are possible; information about lot/batch number can not be obtained.

VAERS ID: 908249 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-12
Onset: 2020-12-12
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Chest discomfort, Hypersensitivity
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Patient has not had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial Patient is not pregnant
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2020506822

Write-up: Chest tightness; Allergy; This is a spontaneous report from a contactable other Health Professional from team and downloaded from the Regulatory Authority-WEB GB-Regulatory Authority-WEBCOVID-20201215082212. Safety Report Unique Identifier GB-Regulatory Authority-ADR 24543246. A 38-years-old female patient received bnt162b2 (BNT162B2, LOT EJ0553), via an unspecified route of administration on 12Dec2020 at single dose for covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. The patient experienced chest tightness and Allergy on 12Dec2020, which Caused/Prolonged Hospitalisation in Dec2020, Other Medically Important Condition. The events resulting in Accident and Emergency (A&E) admission requiring intravenous (IV) hydrocortisone; intramuscular (IM) chlorphenamine and 3 days of fexofenadine. The patient was discharged on 14Dec2020. The event outcome of chest tightness was unknown; outcome of allergy was recovered with sequela on 14Dec2020. The patient did not have symptoms associated with COVID-19, had not been tested/or has had an inconclusive test for COVID-19. No follow-up attempts possible. No further information is expected.

VAERS ID: 909130 (history)
Form: Version 2.0
Age: 51.0
Sex: Male
Location: New York
Vaccinated: 2020-12-16
Onset: 2020-12-20
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2020-12-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Acute myocardial infarction, Catheterisation cardiac abnormal, Coronary artery occlusion, Troponin increased
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Bupropion
Current Illness: None
Preexisting Conditions: Psoriasis, psoriatic arthritis, seasonal affective disorder
Allergies: Amoxicillin

Diagnostic Lab Data: Cardiac cath with 100% RCA occlusion and 75% LAD occlusion, elevated troponin
CDC Split Type:

Write-up: Acute NSTEMI with symptom onset 4 days after vaccination

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VAERS ID: 909146 (history)
Form: Version 2.0
Age: 46.0
Sex: Male
Location: Texas
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / SC

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Disorientation, Dyspnoea, Heart rate increased, Hyperhidrosis, Hypertension, Paraesthesia
SMQs:, Anaphylactic reaction (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hypertension (narrow), Cardiomyopathy (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: I developed tingling all over body with disorientation, rise in heart rate to 140 mint/mints with New onset of severe HTN into 190 mmHg and diaphoresis, followed by SOB and Chest pain. I required IV solucotef, IV tendril and pepcid. Then i
Current Illness: Migraine
Preexisting Conditions: Migraine
Allergies: Levaquin-zithromax
Diagnostic Lab Data: see listed before

CDC Split Type:

Write-up: listed before

VAERS ID: 909379 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-26

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Cardiac monitoring, Chest X-ray, Computerised tomogram abdomen, Computerised tomogram pelvis, Computerised tomogram thorax, Electrocardiogram, Hypotension, Laboratory test, Nausea, Presyncope, Tachycardia, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations: mild near syncope at age 18 (2010) after Gardasil dose
Other Medications: bupropion PO 300mg daily fexofenadine PO 180mg daily Mirena IUD
Current Illness:
Preexisting Conditions:
Allergies: sulfa drugs, mango skin
Diagnostic Lab Data: 12/23/20 emergency department lab workup, chest x-ray, chest/abd/pelvis CT scan, EKG x3, inpatient hospital labs, cardiac monitoring
CDC Split Type:

Write-up: near syncope, hypotension, nausea/vomiting, tachycardia (120-150) within 5 minutes of administration. did not resolve and worsened within 1 hour. Pt went to ER for workups. Received IV benadryl without improvement. Admitted to hospital overnight for

continuous cardiac monitoring. Improved overnight and discharged in the afternoon 12/24/20.

VAERS ID: 909490 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: Washington
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-27

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaemia, Dizziness, Haemoglobin decreased, Nausea, Palpitations, Sinus tachycardia
SMQs:, Acute pancreatitis (broad), Haematopoietic erythropenia (broad), Haemorrhage laboratory terms (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none
Preexisting Conditions: none
Allergies: pollen
Diagnostic Lab Data: Anemia - Hb 9.5 tachycardia 120s
CDC Split Type:

Write-up: The patient developed palpitations, lightheadedness and nausea and came to the ED and was found to have sinus tachycardia. Unclear if it is a vaccine reaction or due to anxiety or severe iron deficiency anemia.

VAERS ID: 909577 (history)

Form: Version 2.0
Age: 60.0
Sex: Female
Location: Michigan
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-27

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Dyspnoea, Full blood count, Laboratory test, Swelling
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Vestibular disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Estrogen, Flonase, Protonix, Glucosamine, Albuterol
Current Illness: Viral respiratory infection
Preexisting Conditions: Allergic rhinitis, GERD, endometriosis
Allergies: Avocado, Banana, Latex, Neosporin, Nystatin, Sulfa, Pineapple
Diagnostic Lab Data: Routine chemistries, CBC
CDC Split Type:

Write-up: Dizziness, dyspnea, neck swelling

VAERS ID: 909586 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-17
Onset: 2020-12-26
Days after vaccination: 9

Submitted: 0000-00-00
Entered: 2020-12-27

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: COVID-19, Chest X-ray abnormal, Exposure to SARS-CoV-2, Pneumonia
SMQs:, Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness: Unknown
Preexisting Conditions: Unknown
Allergies: Unknown
Diagnostic Lab Data: Chest exam and e-x-ray"s (hospitalized)
CDC Split Type:

Write-up: The patient was vaccinated on 12/17/20. Wife was diagnosis with COVID-19 on 12/18/20. He was diagnosis with COVID-19 on 12/21/20. Symptoms woresen on 12/26/20. And he had chest exam (x-ray"s), pneumonia bi-lateral and he was hostipalized on 12/26/20.

VAERS ID: 909635 (history)
Form: Version 2.0
Age: 46.0
Sex: Male
Location: Maryland
Vaccinated: 2020-12-21
Onset: 2020-12-22
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-27

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Other **Purchased by:** ?

Symptoms: Atrial fibrillation, Cardioversion, Chest discomfort, Dyspnoea, Echocardiogram normal, Palpitations, Presyncope
SMQs:, Anaphylactic reaction (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: metformin, Jardiance, Losartan, HCTZ
Current Illness:
Preexisting Conditions: DM, HTN
Allergies: Amoxicillin
Diagnostic Lab Data: Normal echocardiogram
CDC Split Type:

Write-up: Palpitations, shortness of breath, chest tightness, presyncope, which led to New onset atrial fibrillation with rapid ventricular response and required synchronized cardioversion and hospitalization. Discharged on anticoagulation and beta-blocker.

VAERS ID: 909720 (history)
Form: Version 2.0
Age: 41.0
Sex: Male
Location: Minnesota
Vaccinated: 2020-12-22
Onset: 2020-12-23
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-27

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain lower, Appendicectomy, Appendicitis, Computerised tomogram abnormal, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome

(broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: 12/24- CT revealed appendicitis, appendectomy performed
CDC Split Type:

Write-up: 12/23- began to experience intermittent right lower quadrant pain in the morning, fever of 100.4 F in the evening which subsided with ibuprofen. 12/24- no fever noted but intermittent right lower quadrant pain continued, seen at the Health Clinic, sent to Hospital ER for CT scan, diagnosed with appendicitis, appendectomy performed.

VAERS ID: 909778 (history)
Form: Version 2.0
Age: 20.0
Sex: Female
Location: Virginia
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-27

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Dysphagia, Electroencephalogram, Pruritus, Psychogenic seizure, Swollen tongue, Tongue pruritus, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Anticholinergic syndrome (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Depovera
Current Illness:
Preexisting Conditions: Asthma
Allergies: Morphine, penicillin, Shellfish, iodine, honey, pollen, dust
Diagnostic Lab Data: EEG
CDC Split Type:

Write-up: 8:27 Hives, itching all over chest and on tongue, tongue swelling, slight difficulty swallowing. 8:30 epi administered/ benedrly 8:32 pseudo seizures begin off and on till the 27th

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VAERS ID: 909911 (history)
Form: Version 2.0
Age: 65.0
Sex: Male
Location: Alaska
Vaccinated: 2020-12-18
Onset: 2020-12-20
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Atrial fibrillation, Cardioversion, Echocardiogram, Electrocardiogram, Laboratory test
SMQs:, Supraventricular tachyarrhythmias (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: HYDROCHLORIATHIAZIDE 25 MG 1/DAY, ALODAPRINE 5MG 1/DAY
Current Illness: HYPERTENSION
Preexisting Conditions: HYPERTENSION
Allergies: N/A
Diagnostic Lab Data: EKG ECHOCARDIOGRAM "VARIETY LABS"
CDC Split Type: vsafe

Write-up: 12/20/2020 12:00 PM DEVELOPED LAPID ATRIAL FIB. WENT TO ER WHEN IT PERSISTED, MEDICAL CENTER. ADMITTED TO INPATIENT; 12/21/2020 - ELECTRICAL CARDIOVERSION. MONITORED OVERNIGHT. 12/22/2020 - DISCHARGED *ON CHRONIC BLOOD THINNER

VAERS ID: 910053 (history)
Form: Version 2.0
Age: 31.0
Sex: Male
Location: Idaho
Vaccinated: 2020-12-24
Onset: 2020-12-24
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Electrocardiogram, Nausea, Pharyngeal swelling, Respiratory viral panel, SARS-CoV-2 test negative, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Amphetamine-dextroamphetamine Zolpidem

Current Illness: Acid Reflux Hiatal Hernia
Preexisting Conditions: Acid Reflux Hiatal Hernia
Allergies: no known
Diagnostic Lab Data: COVID-19 test - negative Respiratory panel - negative CMC EKG
CDC Split Type:

Write-up: 15 minute post vaccination observation patient denied any symptoms. Later in the day patient experienced significant nausea and vomiting followed by mild SOB and throat swelling.

VAERS ID: 910059 (history)
Form: Version 2.0
Age: 37.0
Sex: Female
Location: Kentucky
Vaccinated: 2020-12-23
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Acute kidney injury, Endotracheal intubation, Gait inability, Hypoaesthesia, Pyrexia, Respiratory failure
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Anaphylactic reaction (broad), Angioedema (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Dystonia (broad), Acute central respiratory depression (narrow), Guillain-Barre syndrome (broad), Hypersensitivity (broad), Tumour lysis syndrome (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), Hypokalaemia (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Cetirezen 10 mg Daily, Lasix 20 Mg daily, Gabapentin 400 bid, Glipizide 5 mg daily, lisnopril 5 mg daily, Levothyroxine25mcg daily, Metoxican 7.5 daily, Sertraline 50 daily
Current Illness: none

Preexisting Conditions: Depression, Diabetes ,Hypertension, Hypothyroidism, Obesity
Allergies: Pencillian
Diagnostic Lab Data:
CDC Split Type:

Write-up: Numbness in sole of feet. Unable to walk, develop high fever, resp failure resulting in intubation, acute kidney injury

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VAERS ID: 910133 (history)
Form: Version 2.0
Age: 64.0
Sex: Female
Location: South Dakota
Vaccinated: 2020-12-23
Onset: 2020-12-24
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: C-reactive protein increased, Full blood count normal, Speech disorder, Swollen tongue, X-ray normal
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Dementia (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Guaifenesin, Albuterol Inh, levothyroxine, amlodipine, coreg, Claritin.
Current Illness: none
Preexisting Conditions: Hypothyroidism, Hypertension, menopause
Allergies: environmental (sneezy/watery eyes), ibuprofen,
Diagnostic Lab Data: Labs completed at 1352. CBC WNL. CRP 7.4. Soft tissue neck xray at 1527 with no acute findings.
CDC Split Type:

Write-up: Noted tongue starting to swell on 12/24 at 1030. Started on left side, then progressed to right side. No SOB, difficulty swallowing or breathing, but staff noted difficulty understanding her speech. Presented to ED at 1300. 50mg Benadryl given IV on 12/24 at 1328 and 125mg solumedrol given IV at 1327. Pt reported improvement in tongue swelling at 1630.

VAERS ID: 910202 (history)
Form: Version 2.0
Age: 39.0
Sex: Male
Location: New Jersey
Vaccinated: 2020-12-18
Onset: 2020-12-23
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Arthralgia, COVID-19, Dizziness, Loss of consciousness, Occupational exposure to SARS-CoV-2, SARS-CoV-2 test positive, Seizure, Syncope
SMQs:, Torsade de pointes/QT prolongation (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Arthritis (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: no
Current Illness: no
Preexisting Conditions: no
Allergies: no
Diagnostic Lab Data: Yes/ medical records.
CDC Split Type:

Write-up: Pediatrician working in the hospital. Was exposed the an office contact wo had covid. Shoulder in soreness. At work on Wednesday. Felt lightheaded had to sit in chair. That"s all he reminders. He workup to a CODE team putting oxygen on him. He has a seizure. Took the COVID test has COVID. Admitted to hospital for 2 days. Likely a syncopal event.

VAERS ID: 910251 (history)
Form: Version 2.0
Age: 41.0
Sex: Male
Location: Kentucky
Vaccinated: 2020-12-17
Onset: 2020-12-27
Days after vaccination: 10
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Blood magnesium, Blood thyroid stimulating hormone, C-reactive protein, Chest X-ray, Computerised tomogram head normal, Electrocardiogram, Full blood count, Influenza virus test, Lumbar puncture, Metabolic function test, Mobility decreased, Muscular weakness, SARS-CoV-2 test, Urine analysis
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Tendinopathies and ligament disorders (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: amlodipine, nicotine patches, Descovy, azelastine, cetirizine, ibuprofen
Current Illness: none
Preexisting Conditions: hypertension, irritable bowel syndrome, kidney stones, HIV exposure 2 months ago
Allergies: none
Diagnostic Lab Data: 12/28/20 Lumbar puncture, urine analysis, COVID test, inflenza tesk, CBC, CT head was negative, CXR, ECG, TSH, Magnesium, CMP, CRP
CDC Split Type:

Write-up: 41-year-old male who presents to the ED today with a complaint of weakness in his bilateral arms and legs. He states he fell slightly weak yesterday but this morning when he

woke up around 6 AM he was not able to get out of bed because he was so weak. He states he feels like he has no muscle strength in his arms and legs. He denies any fever. He denies any cough or shortness of breath. He denies any chest pain or abdominal pain. He denies any nausea, vomiting or diarrhea. He denies any numbness in his extremities. He denies any neck or back pain. He did receive the first Covid vaccination on December 17.

VAERS ID: 910316 (history)
Form: Version 2.0
Age: 22.0
Sex: Male
Location: Wisconsin
Vaccinated: 2020-12-17
Onset: 2020-12-21
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / OT

Administered by: Private **Purchased by:** ?
Symptoms: Laboratory test, Platelet transfusion, SARS-CoV-2 test negative, Thrombotic thrombocytopenic purpura
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Embolic and thrombotic events, arterial (narrow), Renovascular disorders (broad), Immune-mediated/autoimmune disorders (narrow), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Date: 20201221; Test Name: work-up; Result Unstructured Data: Test Result:unknown results; Test Date: 20201221; Test Name: covid_test; Test Result: Negative
CDC Split Type: USPFIZER INC2020505215

Write-up: TTP; This is a spontaneous report from a non-contactable pharmacist. A 22-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA

VACCINE), intramuscular on 17Dec2020 as a single dose for COVID-19 immunization. The patient did not have any known relevant medical history. The patient had no allergies to medications, food or other products. Prior to the vaccination, the patient was not diagnosed with COVID-19. The patient"s concomitant medications were not reported. It was unknown if the patient received any other vaccines within four weeks prior to the vaccination. On 21Dec2020, the patient experienced thrombotic thrombocytopenic purpura (TTP); which was serious for hospitalization. The clinical course was as follows: The patient went to the emergency room/urgent care and was admitted in the early morning of 21Dec2020 due to TTP. Work-up was ongoing with no known results. On 21Dec2020, the patient also had a COVID-19 test which was negative. The patient was treated with unspecified corticosteroids and platelets. The clinical outcome of the TTP was unknown. The reporter assessed that it was unknown if the TTP was related to the vaccination. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.; Sender"s Comments: Current limited information does not allow a full medically meaningful assessment, especially lack of medical history, concomitant medications, concurrent illness and diagnostic workups such as coagulation test, Combs test, bacterial/virologic/immunological biomarkers to identify the etiology. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 910602 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: California
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Abdominal pain lower, Pancreatitis
SMQs:, Acute pancreatitis (narrow), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes

Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ACETAMINOPHEN; ; SERTRALINE HCL; ZOLOFT
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020506207

Write-up: pancreatitis; acute lower abdominal pain; This is a spontaneous report from a contactable pharmacist. A 46-year-old non-pregnant female patient received the first dose of the bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot Number: EJ1685), intramuscularly on 18Dec2020 at 08:00 (as reported) at 46-years-old at a single dose for COVID-19 immunisation. The patient medical history was not reported. The patient had no known drug allergy (NKDA). Concomitant medications included acetaminophen (MANUFACTURER UNKNOWN), propranolol (MANUFACTURER UNKNOWN), sertraline hcl (MANUFACTURER UNKNOWN), sertraline hydrochloride (ZOLOFT); all taken for an unspecified indication from an unspecified date to an unspecified date (which were received within two weeks of vaccination). On 18Dec2020 at 17:00, the patient experienced pancreatitis and acute lower abdominal pain; which required hospitalization and were assessed as medically significant. The patient was hospitalized for pancreatitis and acute lower abdominal pain for 3 days on unspecified dates. The clinical course was reported as follows: The patient received the vaccine " at some point in the AM on 18Dec2020 (as reported)." That evening, the patient presented to the emergency department (ED) with acute lower abdominal pain. The patient was diagnosed with pancreatitis and was admitted overnight. It was unknown if the patient received any other vaccines within four weeks prior to the COVID vaccine. Prior to the vaccination, it was unknown if the patient was diagnosed with COVID-19. Since the vaccination, the patient had not been tested for COVID-19. Therapeutic measures were taken as a result of pancreatitis and acute lower abdominal pain. The clinical outcome of the events was recovering.; Sender"s Comments: The information currently provided is too limited to make a meaningful medical assessment. Other than a temporal association , there is no evidence or argument to suggest a causal relationship between BNT162B2 and the events pancreatitis and acute lower abdominal pain. The events are likely due to an underlying medical condition. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 910624 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot /	Site / Route

	Dose	
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Basal ganglia haemorrhage, Condition aggravated, Hypertension
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Neuroleptic malignant syndrome (broad), Haemorrhagic central nervous system vascular conditions (narrow), Hypertension (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Hypertension
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020507193

Write-up: Admitted to the hospital with hypertensive basal ganglia bleed, had a head bleed; This is a spontaneous report from a contactable consumer. A male patient of unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration in 2020 at single dose for COVID-19 immunization. Medical history included hypertension. The patient"s concomitant medications were not reported. After vaccination, the patient was admitted to the hospital with hypertensive basal ganglia bleed, had a head bleed in 2020. The outcome of event was unknown. Information on the lot/batch number has been requested.

VAERS ID: 910626 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 2020-12-16
Onset: 2020-12-19
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) /	- / 1	RA / -

PFIZER/BIONTECH		

Administered by: Other **Purchased by:** ?
Symptoms: Respiratory distress, SARS-CoV-2 test negative
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Hypersensitivity (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201219; Test Name: covid test/Nasal Swab; Test Result: Negative
CDC Split Type: USPFIZER INC2020507536

Write-up: resp distress; This is a spontaneous report from a non-contactable consumer (patient). An elderly male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration at 16Dec2020 12:00 pm at single dose for covid-19 immunization. Vaccine location was right arm and it was the first dose. The patient medical history and concomitant medications were not reported. No known allergies. Patient was not diagnosed with COVID-19 prior to vaccination. The patient didn''t receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient experienced respiratory distress on 19Dec2020, he was hospitalized for three days. Patient received treatment for the adverse event. Since the vaccination, the patient has been tested for COVID-19 with nasal swab on 19Dec2020, it was negative. The action taken in response to the events for BNT162B2 was not applicable. The outcome of events was unknown. The event was serious, the seriousness criteria was Caused/prolonged hospitalization. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 910641 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: Maryland
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac monitoring, Chest X-ray, Computerised tomogram, Dizziness, Emotional distress, Fatigue, Feeling abnormal, Heart rate decreased, Hypersomnia, Injection site pain, Muscular weakness, Pain in extremity, Palpitations, Paraesthesia, Pulse abnormal, Scan with contrast, Ventricular extrasystoles, Vertigo
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Ventricular tachyarrhythmias (narrow), Dementia (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Extravasation events (injections, infusions and implants) (broad), Cardiomyopathy (broad), Depression (excl suicide and self injury) (broad), Vestibular disorders (narrow), Tendinopathies and ligament disorders (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Date: 20201222; Test Name: Blood pressure; Result Unstructured Data: Test Result:163/76; Comments: Normal base line is 130s/80s maybe lower; Test Date: 20201222; Test Name: cardiac monitor; Result Unstructured Data: Test Result:Results are pending; Test Date: 20201222; Test Name: Chest X-ray; Result Unstructured Data: Test Result:Unknown result; Comments: Result: Pending; Test Date: 20201222; Test Name: CT scan; Result Unstructured Data: Test Result:Unknown result; Comments: Result: Pending; Test Date: 20201222; Test Name: Heart rate; Result Unstructured Data: Test Result:80s-90s; Comments: 90s and then drop to 48; Test Date: 20201219; Test Name: pulse; Result Unstructured Data: Test Result:Thready; Test Date: 20201222; Test Name: Pulse oximetry; Result Unstructured Data: Test Result:97-99 %
CDC Split Type: USPFIZER INC2020508928

Write-up: Weakness and tingling down left arm; Weakness and tingling down left arm; Lightheaded; PVC''s every 3 beats; emotional too and just very tired; Can not read the vaccination card as she does not have her glasses; Palpitations; Fatigue; Slept a lot; Thready pulse and vertigo; Thready pulse and vertigo; Soreness in left arm at the injection site and down the left arm; Soreness in left arm at the injection site and down the left arm; This is a spontaneous report from a contactable nurse (patient). A 54-year-old female patient received first dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, reason for no lot number of COVID Vaccine: Can not read the vaccination card as she does not have her glasses, Expiry

Date unknown), via an unspecified route of administration in the left arm on 18Dec2020 at single dose for "Work with COVID patients". Medical history included none. There were no concomitant medications. The patient experienced weakness and tingling down left arm (hospitalization) on 22Dec2020, lightheaded (hospitalization) on 22Dec2020, PVC"s every 3 beats (hospitalization) on 22Dec2020, soreness in left arm at the injection site and down the left arm on 18Dec2020, thready pulse and vertigo on 19Dec2020, fatigue on 20Dec2020, slept a lot on 19Dec2020, palpitations on 21Dec2020. Details as follows: Caller says she received the vaccine, she is a nurse. She got the vaccine on Friday, 18Dec2020. She had soreness in her arm and at the injection site on Friday but that was it. On Saturday (19Dec2020) she noticed a thready pulse, but went on with her day with only a little arm pain. Sunday (20Dec2020) she was fatigued and the thready pulse continued. She slept a lot on Saturday (19Dec2020) and Sunday (20Dec2020). Yesterday (21Dec2020) she felt a little better, but had palpitations here and there. This morning (22Dec2020) she went into work, was very lightheaded, had tingling down her left arm, and had palpitations. So she hooked herself up to a monitor. Her pulse ox was between 97-99%. Her heart rate would be in the 90s and then drop to 48, so she went down to the ED. She has had a CT, and she is throwing PVC"s every 3 beets. She has not been admitted as they are still waiting for results. She is still in the ED. They did a CT to see if there was a possible clot. On 18Dec2020 she received the vaccine around 2 PM. She had soreness at the injections site and down the left arm, which went away by Sunday (20Dec2020). She now (22Dec2020) has weakness and tingling down the left arm. It was never red or anything at the injection site. Saturday, 19Dec2020, she had thready pulse and Vertigo which lasted until Sunday 20Dec2020. She would be laying in bed and try to flip to the other side and having vertigo. When the fatigue started on Sunday (20Dec2020) she did not feel like herself. She was very emotional too and just very tired. Since she went to the ED she has had a CT scan, one with contrast and one without. She had a chest X-ray, and she is on a cardiac monitor. Results are pending. She has Trigeminy PVCs. She says she never goes to the hospital. But she is not admitted yet (pending clarification). Can not read the vaccination card as she does not have her glasses. Unable to read off the NDC, lot, and expiration date. History: Has been on the same vitamins for two years with nothing new. Blood pressure: Normal base line is 130s/80s maybe lower. Heart rate: Currently within her normal limits of 80s-90s. Depending on what happens, it was asked if she should get the second dose. The patient underwent other lab tests and procedures which included blood pressure measurement: 163/76 on 22Dec2020, chest x-ray: unknown result on 22Dec2020 (Result: Pending), computerised tomogram (CT scan): unknown result on 22Dec2020 (Result: Pending), heart rate: 80s-90s on 22Dec2020, Pulse oximetry: 97-99 % on 22Dec2020, cardiac monitor: results are pending on 22Dec2020. The outcome of events weakness and tingling down left arm, pvc"s every 3 beats, lightheaded, palpitations and fatigue was not recovered. The outcome of the event soreness in left arm at the injection site and down the left arm was recovered on 20Dec2020. The outcome of the events thready pulse and vertigo was recovered on 20Dec2020. The outcome of the event slept a lot was recovered on 20Dec2020. The outcome of other events was unknown. Information on the lot/Batch number has been requested.; Sender"s Comments: Based on available information, a possible contributory role of the subject vaccine cannot be excluded for the reported events due to temporal relationship. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics, counteractive treatment measures and concomitant medications. This case will be reassessed once additional information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 910723 (history)
Form: Version 2.0
Age: 61.0
Sex: Female
Location: New Jersey
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	MODERNA / UNK	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest X-ray normal, Chest discomfort, Cough, Neutrophil percentage increased, Throat clearing
SMQs:, Anaphylactic reaction (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Albuterol 90 mcg/ Aerosol 2 puffs 4 times day Famotidine 20mg po daily Prednisone 20mg po 2 times day Diphenhydramine 25 mg po Q 6 hrs prn
Current Illness: UTI 11/28/2020
Preexisting Conditions: Hx Asthma Hx Uterine Sarcoma
Allergies: MRI Dye, NSAIDs
Diagnostic Lab Data: 12/26 CXR: Negative, no pneumonia, edema or pneumothorax Lab work: Neutrophils 83.6
CDC Split Type:

Write-up: 10 minutes after the vaccination, she began clearing her throat, within 30 minutes began coughing, which led to chest tightness. Was evaluated in the ER and admitted for observation. Given: Prednisone 40mg po, Benadryl 25mg po Duoneb x 3 and Pepcid 20mg

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VAERS ID: 910750 (history)
Form: Version 2.0
Age: 76.0
Sex: Male

Location: Oregon
Vaccinated: 2020-12-22
Onset: 2020-12-25
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Angiogram cerebral normal, Arteriogram carotid normal, Blood cholesterol normal, Blood creatinine normal, Blood potassium normal, Blood sodium decreased, Blood triglycerides normal, Blood urea normal, Creatinine renal clearance decreased, Drooling, Dysarthria, Haematocrit normal, Haemoglobin normal, High density lipoprotein decreased, Hypertensive urgency, International normalised ratio normal, Magnetic resonance imaging brain normal, Platelet count normal, Posture abnormal, Unresponsive to stimuli, Wheelchair user, White blood cell count normal
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Dyslipidaemia (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Dyskinesia (broad), Dystonia (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hyponatraemia/SIADH (narrow), Hypertension (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Lipodystrophy (broad), Hypotonic-hyporesponsive episode (broad), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Tylenol 325mg - 2 tablets Famotidine 20mg - 1 tablet BID Fenofibrate 160mg - 1 tablet daily Vitamin D 2000iu - 1 tablet daily Hydralazine 25mg - 1 tablet TID Amlodipine Besylate 5mg - 1 tablet BID Atorvastatin 40mg - 1 tablet HS Donepezil
Current Illness: none noted
Preexisting Conditions: hypertension associated with Stage 3 chronic kidney disease due to type II diabetes hemiparesis - R dominant side vascular dementia w/out behavioral disturbance hx of stroke with residual deficit -2005 - R medial thamamus -2018 - L basal ganglia paroxysmal atrial fibrillation type II diabetes dysphagia as late effect of stroke sleep apnea hyperlipidemia
Allergies: Aspirin, Ibuprofen, Losartan, Simvastatin, ACE inhibitors, Penicillins
Diagnostic Lab Data: MRI brain - no acute infarct CTA Head/Neck scan - failed to show a new area of ischemia Labs - no infection noted, WBC unrevealing, creatinine clearance 56.5

ml/min (based on SCr of 1.22mg/dL) - at baseline, sodium 134, potassium 4.8, BUN 13, PT INR 1.1, HGB 17.0, platelets 287, hematocrit 51.0, chol 167, trig 131, HDL 35
CDC Split Type:

Write-up: Patient was found slumped over in wheelchair, drooling and unable to respond/follow simple instructions; sent to ED for evaluation. CT scans and MRI NEGATIVE for new/recent stroke; resident slurred speech likely due to hypertensive urgency

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VAERS ID: 910822 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: Mississippi
Vaccinated: 2020-12-22
Onset: 2020-12-23
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angioedema, Enlarged uvula, Hypersensitivity, Lip oedema, Lip swelling, Nasal congestion, Oedema mouth, Palatal oedema, Paraesthesia oral, Sneezing, Throat tightness, Tongue oedema
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: AMLODIPINE, HYDROCHLOROTHIAZIDE, CAMRESE, ICAR-C, LEXAPRO, ACZONE, RETIN A MICRO PUMP
Current Illness: NONE
Preexisting Conditions: ANEMIA
Allergies: NORCO, SPIRONOLACTONE
Diagnostic Lab Data:
CDC Split Type:

Write-up: SNEEZING, STUFFY NOSE, LIP SWELLING, TINGLING MOUTH, THROAT CLOSING, UVULA SWOLLEN 1. Allergic reaction (Allergy, unspecified, initial encounter) . Pt.

received Moderna COVID vaccine the morning of 12/22/20 and developed angioedema reaction the afternoon of 12/23 Vaccine is only new medication reported by pt. - denies any new products, foods, etc History of hives with spironolactone 2.5 months ago; taking HCTZ now - unknown if reaction is related to this med rather than vaccine? Due to severity of her reaction and uncertainty if all attributable to vaccine versus some other culprit I would recommend Epi pen at time of discharge Has received IM epinephrine, racemic epi neb, Solu-Medrol, Benadryl, Pepcid in the ED Continue with Benadryl 25mg IV q6h, Pepcid 20mg IV q12h, and Solu-Medrol 40mg IV q6h 2. Angioedema (Angioneurotic edema, initial encounter) . Arrived to the ED with R. side lip, tongue, and uvula and oral cavity edema Airway patent, room air O2 sats WNL - monitor throughout the night for any worsening of angioedema or airway involvement Treatment as above

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VAERS ID: 910948 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Erythema
SMQs:, Anaphylactic reaction (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: None known
Preexisting Conditions: Asthma
Allergies: Green beans, cinnamon, shrimp, coconut, egg, iodine, latex, naproxen, oseltamivir, sesame seed, guaifenesin
Diagnostic Lab Data:

CDC Split Type:

Write-up: Difficulty breathing 5 minutes after receiving first dose of Covid-19 vaccine by Pfizer/BioNTech, small erythematous spots to bilateral arms.

VAERS ID: 910959 (history)
Form: Version 2.0
Age: 33.0
Sex: Male
Location: California
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK#5730 / UNK	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Alanine aminotransferase increased, Angiogram cerebral normal, Arteriogram carotid normal, Aspartate aminotransferase increased, Blood albumin normal, Blood alkaline phosphatase normal, Blood bilirubin normal, Blood chloride normal, Blood creatinine normal, Blood glucose normal, Blood potassium decreased, Blood sodium decreased, Blood urea normal, Carbon dioxide decreased, Computerised tomogram head normal, Confusional state, Electroencephalogram normal, Eosinophil count normal, Haematocrit decreased, Haemoglobin decreased, Head discomfort, Hypoaesthesia, Hypokalaemia, Language disorder, Magnetic resonance imaging brain normal, Mean cell haemoglobin concentration normal, Mean cell volume normal, Mean platelet volume decreased, Migraine, Peripheral coldness, Platelet count normal, Red blood cell count decreased, Salivary hypersecretion, Troponin I normal, White blood cell count normal
SMQs:, Liver related investigations, signs and symptoms (narrow), Haematopoietic erythropenia (narrow), Peripheral neuropathy (broad), Haemorrhage laboratory terms (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hyponatraemia/SIADH (narrow), Chronic kidney disease (broad), Myelodysplastic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Hypokalaemia (narrow), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: No
Current Illness: No
Preexisting Conditions: No
Allergies: No known allergies
Diagnostic Lab Data: CMP: Recent Labs 12/23/20 1525 12/23/20 1650 NA 135* -- K 3.3* -- CL 101 -- CO2 20.3 -- GLUCOSE 124 106 BUN 11.0 -- CREATININE 1.1 -- ALBUMIN 3.8 -- ALKPHOS 67 -- ALT 37 -- AST 41* -- BILITOT 0.4 -- ? ? CBC: Recent Labs 12/23/20 1525 WBC 10.00 EOSABS 0.2 RBC 4.51 HGB 13.5 HCT 39.1 MCV 86.7 MCHC 34.6 PLT 343 MPV 7.1* ? Results from last 7 days Lab Units 12/23/20 1525 TROPONIN I ng/mL <0.017 ? Results from last 7 days Lab Units 12/23/20 1525 TROPONIN I ng/mL <0.017? ? Lab Results Component Value Date ? HGB 13.5 12/23/2020 ? ? ? Lab Results Component Value Date ? HGB 13.5 12/23/2020 ? ?
CDC Split Type:

Write-up: 33 y.o.?male?with no significant past medical history except for obesity who has been working as a nurse in the emergency room department in our hospital and today he received COVID-19 vaccine and 30 minutes later patient started having increased saliva, cold hands and feet, left-sided pressure-like headache and some numbness in his legs at the same time he suddenly started talking only in first language and lost his ability to speak in second language. ?He understand second language but replying first language stating that he is talking in second language. ?On exam he was alert oriented confused by people not understanding his second language stating that his numbness and cold feeling in the hands and feet have improved. ?Initially patient received 10 mg of Decadron for possible allergic reaction he had a head CT scan that was negative and his labs were remarkable only for hypokalemia. ?Patient had no prior history of any neurological symptoms he was advised admission to the hospital for observation ? ? Patient symptoms resolved next day,he is alert oriented able to communicate in second language he had a head MRI and head neck MRA that came back negative and had an EEG that showed no seizure activities. Patient was seen in neurology consultation who felt that patient most likely had an episode of migraine headache. Patient is going to be discharged home and to have a follow-up with his primary care physician next week ?

VAERS ID: 910996 (history)
Form: Version 2.0
Age: 64.0
Sex: Female
Location: California
Vaccinated: 2020-12-24
Onset: 2020-12-24
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthma, Chest discomfort, Condition aggravated, Dizziness, Malaise,

Palpitations, Supraventricular tachycardia
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (narrow), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Advair Diskus 500 mcg-50 mcg inhalation powder 1 inhal, inhalation, BID amLODIPine 5 mg oral tablet 5 mg = 1 tab, PO, daily calcium carbonate 1 tab, PO, daily cyclobenzaprine 10 mg oral tablet 10 mg = 1 tab, PRN, PO, QHS Euthyrox 112 mcg (0
Current Illness: Asthma exacerbation, started oral steroids 12/21
Preexisting Conditions: Asthma, hypertension, dyslipidemia, hypothyroidism, appendectomy, history of uterine fibroids status post endometrial ablation
Allergies: Tape-Blisters sulfa drugs-Hives Toradol-Itching Vicodin-Rash
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient reports taking Pfizer Covid vaccine and 2 hours after that she reports feeling not well. She recorded that one of the side effects was heart arrhythmias and hence she had her coworker checked her rhythm and she reported that her heart rate was in 180s and hence she was brought to the emergency department for further evaluation. She reports at the time she had palpitations and felt mild lightheadedness and dizziness. She was found to be in SVT with heart rate in the range of 1 80-220 and she received 1 dose of 6 mg Adenoscan after which she converted to normal sinus rhythm. At the time of my evaluation she is in normal sinus rhythm with heart rate in the range of 90-100. She denies any further palpitations. She reports she had chest tightness for the last 3 days which was assumed to be secondary to asthma and for which she was prescribed prednisone. Currently with the prednisone she does not feel any further chest tightness. She denies any chest pain shortness of breath, fever or chills. She reports remote history of arrhythmia following her foot surgery in the past however does not recall what arrhythmia she had at that time.

VAERS ID:	911025 (history)
Form:	Version 2.0
Age:	43.0
Sex:	Female
Location:	California
Vaccinated:	2020-12-20
Onset:	2020-12-20
Days after vaccination:	0
Submitted:	0000-00-00

Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Chest X-ray abnormal, Chills, Cough, Diarrhoea, Headache, Hypotension, Laboratory test normal, Lung opacity, Myalgia, Occupational exposure to SARS-CoV-2, Pyrexia, SARS-CoV-2 test positive, Tachypnoea
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (narrow), Asthma/bronchospasm (broad), Interstitial lung disease (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Noninfectious diarrhoea (narrow), Respiratory failure (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: amLODIPine 10 mg PO daily ascorbic acid 250 mg PO dailyJardiance 25 mg oral tablet 25 mg = 1 tab, PO, daily Jardiance 25 mg PO daily losartan 100 mg PO daily metformin 1000 mg PO BID Omega-3 1 cap PO daily simvastatin 20 mg oral tablet PO Q
Current Illness: none
Preexisting Conditions: Type 2 diabetes, hypertension
Allergies: Lactose-diarrhea
Diagnostic Lab Data: 12/26 SARS-CoV-2 PCR positive In the emergency department the patient was mildly hypotensive 97/54, low-grade fever 37.4, tachypneic, saturating 90% on room air, labs showed no leukocytosis, normocytic anemia, hyponatremia and hypochloremia, chest x-ray showed bilateral groundglass opacities.
CDC Split Type:

Write-up: Patient presents to the emergency department 12/26 complaining of dry cough associated with fever and chills and headache associated with myalgia and diarrhea for 1 week duration. She had Covid vaccine 12/20 and the symptoms started the same night, she denied any sick contacts at home however she works at the Covid unit and reports constant exposure to sick Covid patients.

VAERS ID: 911035 (history)
Form: Version 2.0
Age: 47.0
Sex: Female

Location: Alabama
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram cerebral normal, Arteriogram carotid normal, Blindness, Cerebrovascular accident, Computerised tomogram head normal, Electrocardiogram normal, Eye irritation, Frustration tolerance decreased, Full blood count normal, Incoherent, Loss of consciousness, Metabolic function test normal, Migraine, Panic reaction, Posture abnormal, Posturing, Seizure, Transient ischaemic attack
SMQs:, Torsade de pointes/QT prolongation (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Convulsions (narrow), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Dystonia (broad), Psychosis and psychotic disorders (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Glaucoma (broad), Optic nerve disorders (broad), Corneal disorders (broad), Retinal disorders (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vitamin D weekly Fluoxetine Daily
Current Illness:
Preexisting Conditions: Depression
Allergies: No known food or medication allergies Seasonal Allergies ~ 5-6 years
Diagnostic Lab Data: CTA- head/ neck- normal CT Scan head unremarkable EKG- sinus rhythm without ischemic changes CBC/CMP fully unremarkable
CDC Split Type:

Write-up: Approximately 7 hours after receiving the vaccine patient who is a L&D Nurse return to work in her area. She describes that after finishing with a C-section she felt burning in both of her eyes (she thought this feels like an allergic reaction, but I am not sweating and haven"t rubbed my eyes). She went to the restroom to get a cloth to wash her eyes; afterwards she reports her vision went totally black in both eyes. She reports feeling frustrated that no one came to help and some panic in trying to figure out how to get out of the restroom.

She did make it out of the bathroom. Her Staff reports she postured and turned arms inward, head going to one side and passed out. They also report ~ 10 minutes of incoherent conversation and stating "I got the vaccine, maybe I was given the wrong thing and now I"m blind". Upon waking, patient vision fully restored and patient does not remember incoherent conversation. Differential diagnosis- TIA vs. CVA $g seizure disorder$ggg complex migraine

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VAERS ID: 910285 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-11-13
Onset: 2020-11-13
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Chest discomfort, Dizziness, Dyspnoea, Headache, Paraesthesia, Rash erythematous, SARS-CoV-2 test
SMQs:, Anaphylactic reaction (narrow), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Patient has not had symptoms associated with COVID-19 Patient is not enrolled in clinical trial
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:Negative COVID-19 test; Comments: negative covid-19 test had not tested positive for COVID-19 since having the vaccine

CDC Split Type: GBPFIZER INC2020505342

Write-up: Tight chest; Light-headed feeling; Red rash; Tingly face around the mouth; Headache; Breathlessness; This is a spontaneous report from a contactable consumer from EVDAS team (EU-EC-10007206161) and downloaded from the Regulatory Authority- GB-MHRA-WEBCOVID-20201214163159, Safety Report Unique Identifier GB-MHRA-ADR 24543179. An adult male patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EJ0553), via an unspecified route of administration on 13Nov2020 at single dose for COVID-19 immunization. The patient medical history was not reported. The patient"s concomitant medications were not reported. The patient experienced tight chest on 13Nov2020 with outcome of recovered on 13Nov2020, light-headed feeling on 13Nov2020 with outcome of recovered on 13Nov2020, red rash on 13Nov2020 with outcome of recovering, tingly face around the mouth on 13Nov2020 with outcome of recovered on 13Nov2020, headache on 13Nov2020 with outcome of recovered on 13Nov2020, breathlessness on 13Nov2020 with outcome of recovered on 13Nov2020. The patient was hospitalization due to all these events. The patient underwent lab tests and procedures which included sars-cov-2 test: negative covid-19 test on unknown date. Patient had not tested positive for COVID-19 since having the vaccine. No follow-up attempts possible. No further information expected.

VAERS ID: 910442 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-10
Onset: 2020-12-11
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Pneumonia
SMQs:, Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19 (severe COVID infection)
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2020509356

Write-up: Acute pneumonia/acute multilobar pneumonia; This is a spontaneous report downloaded from the Regulatory authority[GB-MHRA-EYC 00235529], Safety Report Unique Identifier GB-MHRA-ADR 24543842. This is a report received from the Regulatory Authority). A contactable physician reported that an 86-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot no. and expiry date was not reported), intramuscular on 10Dec2020 at single dose for COVID-19 vaccination. Medical history included Covid-19 on unspecified date in 2020. The patient''s concomitant medications were not reported. The patient experienced acute pneumonia on 11Dec2020. The patient was hospitalized on unspecified date due to the event. Furthermore, it was reported that the patient had acute multilobar pneumonia not responding to antibiotics including co-amoxiclav, clarithromycin and Tazocin. It could be reaction to COVID vaccine day before. The patient had severe COVID infection earlier in year. The outcome of the event was not recovered. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 910446 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-10
Onset: 2020-12-10
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood glucose, Blood pressure measurement, Cold sweat, Dizziness, Headache, Heart rate, Nausea, Pallor, Peripheral coldness, Respiratory rate, Somnolence, Tremor, Tryptase
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Parkinson-like events (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: No previous medical history. Fit and well. Full childhood vaccinations completed with no problems. Unsure if patient has had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial.
Allergies:
Diagnostic Lab Data: Test Date: 20201210; Test Name: Blood glucose; Result Unstructured Data: Test Result:Blood glucose; Comments: 5.3; Test Date: 20201210; Test Name: Blood pressure; Result Unstructured Data: Test Result:Blood pressure; Comments: 165/90; Test Date: 20201210; Test Name: Heart rate; Result Unstructured Data: Test Result:Heart rate; Comments: 95; Test Date: 20201210; Test Name: Respiratory rate; Result Unstructured Data: Test Result:Respiratory rate; Comments: 23; Test Date: 20201210; Test Name: Blood tryptase; Result Unstructured Data: Test Result:Blood tryptase
CDC Split Type: GBPFIZER INC2020509346

Write-up: Headache; Light-headed; Nausea; Drowsiness; Pale; Clammy; Shakiness; Peripheral coldness; his is a spontaneous report downloaded from the Medicines Agency (MA) WEB, GB-MHRA-ADR 24541968 and GB-MHRA-WEBCOVID-20201210134639. This is a report received from the Regulatory Authority. A contactable pharmacist reported that a 24-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, batch/lot number EJ0553), via an unspecified route of administration on 10Dec2020 at single dose for covid-19 immunisation. The patient medical history was reported as, no previous medical history. Fit and well. Full childhood vaccinations completed with no problems. Unsure if patient has had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Concomitant medications were not reported. The patient previously took influenza virus and experienced no adverse event. The patient experienced headache on 10Dec2020, light-headed on 10Dec2020, nausea on 10Dec2020, drowsiness on 10Dec2020, pale on 10Dec2020, clammy on 10Dec2020, shakiness on 10Dec2020, peripheral coldness on 10Dec2020. The events for considered serious for being medically significant and hospitalization. It was reporter that Within minutes of vaccination complained of feeling shaky and lightheaded, looked pale and clammy, peripherally cold. Observations taken blood pressure 165/90, respiratory rate 23, heart rate 95. Unable to obtain sats due to cold fingers. 15 minutes later complained of feeling nauseous and of "banging" headache. Paracetamol 1g taken 30 minutes later and this helped with good effect for headache but continued to feel nauseous. Blood pressure settled, resps and sats normal but felt very sleepy. As still light headed if sat up blood glucose checked which was 5.3. Taken to Emergency Department for further observation. Bloods taken for mast cell tryptase. Indication for influenza vaccine was: patient is a nurse. Patient is not enrolled in clinical trial. Patient is not pregnant. The outcome of the events headache, nausea, drowsiness (somnolence), pale (pallor), clammy (cold sweat), shakiness (tremor) and peripheral coldness were unknown at the time of this report. The outcome of the event light-headed (dizziness) was recovering at the time of this report. No follow-up attempts are possible; information about batch number cannot be obtained. No further information is expected.

VAERS ID: 910449 (history)
Form: Version 2.0
Age: 64.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-10
Onset: 2020-12-10
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ 0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Abdominal pain upper, Dry mouth, Electrocardiogram, Headache, Hot flush, Presyncope
SMQs:, Acute pancreatitis (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Medical history not known. No health conditions raised on pre vaccine assessment. Confirmed no allergies, no anaphylaxis, no EpiPen, no flu vaccine within 7 days, no anti coagulant concerns, no blood disorders, no concerns with previous vaccinations. Patient has not had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial. Patient is not pregnant.
Allergies:
Diagnostic Lab Data: Test Name: ECG; Result Unstructured Data: Test Result:unknown redults
CDC Split Type: GBPFIZER INC2020510015

Write-up: Headache; Hot flush; Dry mouth; Near fainting; Stomach pain; This is a spontaneous report downloaded from the Medicines Agency (MA) WEB report number GB-MHRA-ADR 24542183 and GB-MHRA-WEBCOVID-20201210222743. A contactable other healthcare professional (HCP) reported that a 64-year-old female patient received BNT162B2

(Pfizer-BioNTech COVID-19 vaccine, Batch numer: EJ 0553), via an unspecified route of administration on 10Dec2020 at single dose for covid-19 immunisation. The patient medical history was not reported. No health conditions raised on pre vaccine assessment. Confirmed no allergies, no anaphylaxis, no EpiPen, no flu vaccine within 7 days, no anti coagulant concerns, no blood disorders, no concerns with previous vaccinations. Patient has not had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial. Patient is not pregnant. The patient''s concomitant medications were not reported. On 10Dec2020, patient experienced dry mouth, headache, hot flush, near fainting, stomach pain. Events were serious due to hospitalization. Patient felt above symptoms at around 14:30 (2-3 hours post vaccination). Treated in staff members clinical area, 12 Lead electrocardiogram (ECG), cannulation, transferred to Emergency for support. Emergency plan to discharge for this evening following blood gas. Patient has not tested positive for COVID-19 since having the vaccine. The outcome of events was recovering. No follow-up attempts are possible. No further information is expected.

VAERS ID: 910460 (history)
Form: Version 2.0
Age: 23.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-11
Onset: 2020-12-11
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0533 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Dysgeusia, Dyspnoea, Heart rate, Nausea, Respiratory rate, SARS-CoV-2 test, Throat irritation, Throat tightness, Wheezing
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (broad), Asthma/bronchospasm (broad), Taste and smell disorders (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypersensitivity (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: Salbutamol
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201211; Test Name: Blood pressure; Result Unstructured Data: Test Result:117/63; Test Date: 20201211; Test Name: Blood pressure; Result Unstructured Data: Test Result:97/68; Test Date: 20201211; Test Name: Heart rate; Result Unstructured Data: Test Result:103; Test Date: 20201211; Test Name: Heart rate; Result Unstructured Data: Test Result:107; Test Date: 20201211; Test Name: Respiratory rate; Result Unstructured Data: Test Result:22; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:no - negative
CDC Split Type: GBPFIZER INC2020507239

Write-up: Wheezing; Throat tightness; Itchy throat; Breathlessness; Nausea; Taste metallic; This is a spontaneous report from contactable pharmacists downloaded from the Medicines Agency (MA) Regulatory Authority-WEB [GB-MHRA-WEBCOVID-20201211165855], Safety Report Unique Identifier is GB-MHRA-ADR 24542578. A 23-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot#: EJ0533) at single dose on 11Dec2020 at 14:30 for COVID-19 immunization. Medical history was not reported. There was no allergies. Patient had not had symptoms associated with COVID-19. Patient had not been tested/or had had an inconclusive test for COVID-19. Patient was not enrolled in clinical trial. Concomitant medication included salbutamol taken for asthma. The patient experienced throat tightness on 11Dec2020 at 14:35 with outcome of recovered on 11Dec2020; itchy throat, breathlessness, nausea, taste metallic, all on 11Dec2020 at 14:35 with outcome of recovering; and wheezing on an unspecified date with outcome of recovering. The seriousness criteria for all adverse events were caused/prolonged hospitalization and other medically important condition. COVID-19 virus test was no - negative. At 14:30, vaccine was given. At 14:35 metallic taste, itchy throat, nausea occurred and then patient started to complain of tight throat but settled after a couple of minutes. Observations were at 14:35, heart rate 103, blood pressure 117/63, respiratory rate 22; at 14:45, heart rate 107, blood pressure 97/68. Patient still felt "scratchy" and uncomfortable when swallowing so given intramuscular steroid, chlorphenamine and salbutamol nebuliser. Patient was taken to emergency department for observation. Patient had not tested positive for COVID-19 since having the vaccine. No follow-up attempts are possible. No further information is expected.; Sender"s Comments: There is a plausible time relationship between exposure and onset of the events. Causality cannot be completely excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 910508 (history)
Form: Version 2.0
Age: 87.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-14
Onset: 2020-12-15
Days after vaccination: 1

Submitted: 0000-00-00
Entered: 2020-12-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Computerised tomogram head, Ischaemic stroke
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Non ST segment elevation myocardial infarction (2/12 ago)
Allergies:
Diagnostic Lab Data: Test Name: Head CT; Result Unstructured Data: Test Result:unknown
CDC Split Type: GBPFIZER INC2020509382

Write-up: stroke; This is a spontaneous report from a contactable physician downloaded from the Medicines Agency (MA) -WEB and received via Regulatory Authority GB-MHRA-WEBCOVID-20201216110942, Safety Report Unique Identifier GB-MHRA-ADR 24543802. An 87-year-old female patient received BNT162B2, via an unspecified route of administration on 14Dec2020 at single dose for COVID-19 vaccination. Medical history included acute myocardial infarction from an unknown date and unknown if ongoing (prior to 02Dec2020). Patient has not had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial. Concomitant medication included acetylsalicylic acid (ACETYLSALICYLIC ACID) for Non ST segment elevation myocardial infarction, clopidogrel (CLOPIDOGREL) Non ST segment elevation myocardial infarction. The patient experienced stroke (hospitalization) on 15Dec2020. Given COVID vaccine in community on 14Dec2020 then presented with ischaemic stroke on 15Dec2020. Unclear significance of COVID vaccine and event. Patient has not tested positive for COVID-19 since having the vaccine. The patient underwent lab test included (Head CT) on unspecified date and unspecified outcome. The outcome of the event was not recovered. No follow-up attempts possible. No further information expected. Lot number cannot be obtained.

VAERS ID: 911636 (history)
Form: Version 2.0
Age: 25.0

Sex: Female
Location: Florida
Vaccinated: 2020-12-22
Onset: 2020-12-25
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram head, Erythema, Full blood count, Hypoaesthesia, Magnetic resonance imaging, Metabolic function test, Paraesthesia oral, Swelling face, Urine analysis
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zyrtec daily. Orillissa daily.
Current Illness:
Preexisting Conditions: Endometrosis, migraines, anxiety
Allergies: Sulfa, and IV contrast dye
Diagnostic Lab Data: MRI and CT of the head to rule out CVA. CMP, CBC, and UA.
CDC Split Type:

Write-up: I received my vaccine on 12/22 around noon. I woke up at 4am this morning 12/25 with tingling in my lips and tongue, in addition to the left side of my face being swollen. Not entirely sure what these side effects are from exactly but thought it best to report my experiences I was hospitalized on 12/25 to 12/26 because my reaction did get worse. I was experiencing numbness on the left side of my body and swelling, numbness, and redness on the left side of my face as well. The hospital didn?t really give any explanations but I?m only now have mild swelling and numbness on the left side of my face. Everything else has resolved at this point in time.

VAERS ID: 911829 (history)
Form: Version 2.0
Age: 80.0

Sex: Female
Location: Michigan
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Asthenia, Decreased appetite, Unresponsive to stimuli
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Aspirin Tablet 81 MG Give 1 tablet by mouth in the morning Bisoprolol Fumarate Tablet 5 MG Give 1 tablet by mouth in the morning for HTN Flomax Capsule 0.4 MG (Tamsulosin HCl) Give 1 capsule by mouth in the evening Levothyroxine Sodium Ta
Current Illness: U07.1 COVID-19
Preexisting Conditions: F03.91 UNSPECIFIED DEMENTIA WITH BEHAVIORAL DISTURBANCE R26.9 UNSPECIFIED ABNORMALITIES OF GAIT AND MOBILITY M62.81 MUSCLE WEAKNESS (GENERALIZED) Z74.1 NEED FOR ASSISTANCE WITH PERSONAL CARE N13.9 OBSTRUCTIVE AND REFLUX UROPATHY, UNSPECIFIED N13.2 HYDRONEPHROSIS WITH RENAL AND URETERAL CALCULOUS OBSTRUCTION R45.4 IRRITABILITY AND ANGER F06.34 MOOD DISORDER DUE TO KNOWN PHYSIOLOGICAL CONDITION WITH MIXED FEATURES G93.41 METABOLIC ENCEPHALOPATHY I25.10 ATHEROSCLEROTIC HEART DISEASE OF NATIVE CORONARY ARTERY WITHOUT ANGINA PECTORIS F06.2 PSYCHOTIC DISORDER WITH DELUSIONS DUE TO KNOWN PHYSIOLOGICAL CONDITION F06.0 PSYCHOTIC DISORDER WITH HALLUCINATIONS DUE TO KNOWN PHYSIOLOGICAL CONDITION R26.2 DIFFICULTY IN WALKING, NOT ELSEWHERE CLASSIFIED I10 ESSENTIAL (PRIMARY) HYPERTENSION K21.9 GASTRO-ESOPHAGEAL REFLUX DISEASE WITHOUT ESOPHAGITIS E11.9 TYPE 2 DIABETES MELLITUS WITHOUT COMPLICATIONS E78.5 HYPERLIPIDEMIA, UNSPECIFIED E03.9 HYPOTHYROIDISM, UNSPECIFIED F41.9 ANXIETY DISORDER, UNSPECIFIED E11.40 TYPE 2 DIABETES MELLITUS WITH DIABETIC NEUROPATHY, UNSPECIFIED E66.01 MORBID (SEVERE) OBESITY DUE TO EXCESS CALORIES J30.9 ALLERGIC RHINITIS, UNSPECIFIED F33.9 MAJOR DEPRESSIVE DISORDER, RECURRENT, UNSPECIFIED F43.23 ADJUSTMENT DISORDER WITH MIXED ANXIETY AND DEPRESSED MOOD R48.8 OTHER SYMBOLIC DYSFUNCTIONS
Allergies: Latex

Diagnostic Lab Data: N/A
CDC Split Type:

Write-up: Resident found unresponsive in her room. Note from earlier: Resident appears to be weak today. Resident ate a few bites of dinner before refusing the tray. Writer encouraged fluids. Vitals 123/72 80HR BS 166. Will log for Doctor and continue to monitor. Was sent out 911.

VAERS ID: 912137 (history)
Form: Version 2.0
Age: 32.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Eye swelling, Swelling face
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Trazodone 50mg, Wellbutrin 300mg, lexapro 20mg, AirDuo Respiclick, Albuterol prn
Current Illness:
Preexisting Conditions: asthma, depression, migraines
Allergies: Lobster, hymenoptera
Diagnostic Lab Data:
CDC Split Type:

Write-up: Was given the vaccine and about 5 minutes later started having swelling and my eyes and face. It was watched for a few minutes and was assessed by EMS and taken to the emergency department. I was given epinephrine, Benadryl, Solu-Medrol, Pepcid, IV fluids, DuoNebs and observed overnight. I was given multiple rounds of Benadryl, steroids, Pepcid,

DuoNebs

VAERS ID: 912271 (history)
Form: Version 2.0
Age: 50.0
Sex: Male
Location: Louisiana
Vaccinated: 2020-12-16
Onset: 2020-12-17
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Intensive care, Pyrexia
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Subject received vaccination Wednesday Dec 16th in the afternoon. He became symptomatic (shortness of breath, low grade fever) the next day. Went to the Emergency room on Saturday Dec. 26th, 2020 due to shortness of breath, had an 02 Sat of 60%, and was hospitalized in the ICU at another hospital (due to bed unavailability).

VAERS ID: 912574 (history)

Form: Version 2.0
Age: 97.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Public **Purchased by:** ?
Symptoms: Cardiac arrest, Endotracheal intubation, Intensive care
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Angioedema (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Cardiomyopathy (broad), Respiratory failure (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: COVID-19 positive on 12/4/2020. Was still symptomatic at last check
Preexisting Conditions: Hypertension
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Rushed to ER. Has now been tubed and put into the ICU and has had full-cardiac arrest less than 24 hours after receiving the vaccine.

VAERS ID: 912602 (history)
Form: Version 2.0
Age: 83.0
Sex: Male
Location: Ohio
Vaccinated: 2020-12-28
Onset: 2020-12-29

Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Endotracheal intubation, Intensive care
SMQs:, Angioedema (broad), Respiratory failure (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: COVID-19 positive 11/30 with fever, sore throat, dyspnea and difficulty breathing
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Hospitalized 12/29, has now been tubed and put into the ICU

VAERS ID: 912609 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Indiana
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Cardiac monitoring abnormal, Dizziness, Flushing, Heart rate increased,

Supraventricular tachycardia, Tachycardia
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypersensitivity (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient observed for 15 minutes in the clinic after vaccine with no issues. Patient is a NP and left clinic. 10 min after leaving, was in physician lounge and had tachycardia, dizziness, flushing. Hospitalist in lounge recorded pulse as 180 normal rhythm. Patient taken to ED. In SVT - heart rate eventually reduced. Patient released on cardiac monitor to home. Later that night, patient had increased heart rate again - 160"s - while in bed. Admitted to hospital.

VAERS ID: 912669 (history)
Form: Version 2.0
Age: 58.0
Sex: Female
Location: Georgia
Vaccinated: 2020-12-22
Onset: 2020-12-23
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Catheterisation cardiac normal, Chest pain, Computerised tomogram normal, Dyspnoea, Electrocardiogram normal, Troponin normal
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic

procedures (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? Yes
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Migraines, Arthiritis
Allergies: Flexril
Diagnostic Lab Data: EKG, Troponins, CT, Cardiac Cath - all negative, 12/23/2020
CDC Split Type:

Write-up: Chest pain, short of breath. Morphine, sublingual Nitro, IV Nitro drip & Heparin drip

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VAERS ID: 912701 (history)
Form: Version 2.0
Age: 82.0
Sex: Male
Location: Arizona
Vaccinated: 2020-12-18
Onset: 2020-12-23
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730*CSU-C19 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram abnormal, Aspartate aminotransferase increased, Blood albumin increased, Blood alkaline phosphatase increased, Blood creatinine increased, Blood glucose increased, Blood urea increased, COVID-19 pneumonia, Carbon dioxide decreased, Cardiopulmonary failure, Cognitive disorder, Dyspnoea, Eosinophil count decreased, Glomerular filtration rate decreased, Hypernatraemia, Hypoxia, Lymphocyte count decreased, Neutrophil count increased, Protein total increased
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Cardiac failure (narrow), Liver related investigations, signs and symptoms (narrow), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Haematopoietic leukopenia (narrow), Hyperglycaemia/new onset diabetes mellitus (narrow), Systemic lupus erythematosus (broad), Retroperitoneal fibrosis (broad), Dementia (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Acute central respiratory depression (broad), Biliary system related investigations, signs and symptoms (broad), Pulmonary

hypertension (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Dehydration (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: - glipizide 2.5 mg/d - metformin ER 2000 mg/d - insulin sliding scale - lisinopril 10 mg/d - amlodipine 2.5 mg/d - furosemide 20 mg/d - Na Bicarb 650 mg TID added 5/23 by nephrology - atorvastatin 20 mg/d - ASA 81 mg/d - levoth
Current Illness:
Preexisting Conditions: CKD 3 HNT Dementia DMII Hypothyroidism
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: Hospitalization 12/26 for Covid PNA

VAERS ID: 912713 (history)
Form: Version 2.0
Age: 70.0
Sex: Female
Location: California
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / UNK	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Ataxia, Consciousness fluctuating, Dizziness, Palpitations
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Vestibular disorders (broad),

Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Hydrocodone/acetaminophen (nausea/vomiting)
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient reported dizziness, weakness, in and out of consciousness, palpitation and ataxia approximately 10 minutes after vaccination. Sent to ED and diagnosed with potential cerebral mass. Palpitations has occurred more frequently in the last 4 months when she ran out of metoprolol for the treatment of SVT since her PMH retired and she has not established new care yet. Dizziness, palpitations and ataxia resolved by next AM. Follow up with outpatient Neuro and cardiology. Given prescription for metoprolol XL 75 daily.

VAERS ID: 912766 (history)
Form: Version 2.0
Age: 27.0
Sex: Male
Location: Kentucky
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Arrhythmia, Atrial fibrillation, Cardiac flutter, Cardiac telemetry, Dizziness, Hyperhidrosis, Nausea
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Supraventricular tachyarrhythmias (narrow), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Cardiac arrhythmia terms, nonspecific (narrow), Tachyarrhythmia terms, nonspecific (narrow), Vestibular disorders (broad), Hypoglycaemia (broad)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: OTC zinc, magnesium, and MVI
Current Illness: none
Preexisting Conditions: none
Allergies: none
Diagnostic Lab Data: The client reports that he was hooked to telemetry. Any other tests/labs are unknown.
CDC Split Type:

Write-up: Client was vaccinated approx 10:00am at the Health Dept. Client left after 15min without incident. Client went to report to work and began feeling nauseated, dizzy, and feeling as if was going to faint. Client returned to Health Department approx 12:30. Client was diaphoretic, HR was 140bpm, and BP was 178/120. Client was given 50mg/20mL of diphenhydramine PO and EMS was called and client taken to local ER. Client remained A+Ox3 at time of EMS transport. Spoke with client evening of 12/28/2020. He reports he was transferred to another facility and was told was in a-fib. He reports that he is feeling much better but heart still fluttering. Spoke with client afternoon of 12/29/2020. He reports that he is being discharged in a few hours and is being sent home on metoprolol and a baby ASA. He reports they had been working on getting his BP down. He reports that he is feeling better and is going to be scheduled for a heart cath outpatient. He reports that he was initially in a-fib, then a-flutter, and then arrhythmia. He reports that he has been seen by Dr. a Cardiologist, while hospitalized.

VAERS ID: 912834 (history)
Form: Version 2.0
Age: 24.0
Sex: Female
Location: Virginia
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / SYR

Administered by: Other **Purchased by:** ?
Symptoms: Electrocardiogram abnormal, Heart rate increased, Hypoaesthesia, Miosis, Mydriasis, Neurological examination, Peripheral coldness, Pyrexia, Tachyarrhythmia
SMQs:, Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Tachyarrhythmia terms,

nonspecific (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), Hypokalaemia (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Amoxicillin
Diagnostic Lab Data: EKG Showed heart was tachycardic and irregular. Raised fever-99.5. Neurological test to explain dilated and constricted pupils.
CDC Split Type:

Write-up: With five minutes of vaccine Increased heart rate shot up to 198 bpm. Regular resting heart rate for me is in the 50s. Left pupil dilated completely, right people constricted to pin size. Extremities went numb and ice cold. Went to emergency room Per request of EMTs at vaccination center. Was given high-dose Benadryl and Saline IV. Heart rate return to normal and all other symptoms disappeared after treatment.

VAERS ID: 912883 (history)
Form: Version 2.0
Age: 41.0
Sex: Male
Location: Missouri
Vaccinated: 2020-12-01
Onset: 2020-12-23
Days after vaccination: 22
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Joint stiffness
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: -Seroquel 1-300 mg -Gabapentin 2-300 mg -Depakote 3-250 mg -Escitalopram (Lexapro) 10 mg -Copaxone 1-40 mg MWF Vitamin B6 1-100 mg Vitamin D3 3-1000 IU
Current Illness:
Preexisting Conditions: Multiple Sclerosis, Bipolar
Allergies: Zyprexa, Trazadone
Diagnostic Lab Data:
CDC Split Type:

Write-up: Difficulty catching my breath, stiff joints in legs

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VAERS ID: 912954 (history)
Form: Version 2.0
Age: 102.0
Sex: Female
Location: North Dakota
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Activated partial thromboplastin time normal, Adjusted calcium, Alanine aminotransferase normal, Anion gap normal, Aortic valve calcification, Arteriosclerosis coronary artery, Aspartate aminotransferase normal, Atelectasis, Atrial enlargement, Basophil count normal, Basophil percentage, Blood albumin normal, Blood alkaline phosphatase normal, Blood bilirubin increased, Blood calcium, Blood chloride normal, Blood creatinine normal, Blood glucose normal, Blood lactic acid, Blood potassium normal, Blood sodium normal, Blood urea nitrogen/creatinine ratio, Blood urea normal, Brain natriuretic peptide increased, C-reactive protein increased, Carbon dioxide decreased, Cardiac failure congestive, Cardioactive drug level, Cardiomegaly, Computerised tomogram thorax abnormal, Cough, Eosinophil count normal, Eosinophil percentage, Fibrin D dimer, Fluid overload, Glomerular filtration rate normal, Haematocrit decreased, Haemoglobin decreased, International normalised ratio increased, Intervertebral disc degeneration, Lymphocyte count decreased, Lymphocyte percentage decreased, Mean cell haemoglobin concentration normal, Mean cell haemoglobin increased, Mean cell volume increased, Mean platelet volume increased, Monocyte count normal, Monocyte percentage, Neutrophil count normal, Neutrophil percentage increased, Osteoarthritis, Platelet count normal, Pleural effusion, Protein total normal, Prothrombin time prolonged, Pulmonary oedema, Pyrexia, Red blood cell count decreased, Red cell distribution width increased, SARS-CoV-2 test positive, Serum ferritin increased, Spinal disorder, Staphylococcus test negative, Tachycardia, Tachypnoea,

Troponin, White blood cell count normal
SMQs:, Cardiac failure (narrow), Liver related investigations, signs and symptoms (narrow), Liver-related coagulation and bleeding disturbances (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Asthma/bronchospasm (broad), Haematopoietic erythropenia (narrow), Haematopoietic leukopenia (narrow), Haemorrhage laboratory terms (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Biliary system related investigations, signs and symptoms (narrow), Pulmonary hypertension (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Other ischaemic heart disease (narrow), Chronic kidney disease (broad), Arthritis (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (narrow), Infective pneumonia (broad), Dehydration (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: tylenol aspirin Atorvastatin B12 Folate Diltiazem Docusate Digoxin Furosemide Potassium Chloride Levothyroxine Lisinopril Protonix Vitamin D3 Warfarin Polyethylene Glycol
Current Illness: COVID-19 (Diagnosed 10/26/20)
Preexisting Conditions: Diabetes Type II Atrial Fibrillation Hypothyroidism CHF Hypertension Osteoporosis Anemia Cerebrovascular disease Aphasia
Allergies: ibuprofen Lactose
Diagnostic Lab Data: Troponin First 0.071, Second 0.054 BNP-398 Digoxin 1.2 SARS -COV-2 Detected MRSA not detected CTA- FINDINGS: Mild motion artifact somewhat limits the evaluation. No definite pulmonary emboli. ?Mild to moderate diffuse interstitial prominence suggesting edema. ?Small bilateral effusions with mild probable atelectasis in the bases. ? Mild to moderate cardiomegaly with biatrial enlargement. Aortic valvular and coronary artery calcification. ?No definite pathologic lymphadenopathy. Mild to moderate spinal curvature. ? Degenerative changes of the spine and both shoulders. ? IMPRESSION: 1. ?Mild to moderate cardiomegaly with biatrial enlargement. ?Mild to moderate diffuse interstitial prominence suggesting edema with small bilateral effusions. ?Findings suggest an element of fluid overload/congestive failure. 2. ?Additional nonacute/chronic findings, as detailed above. 3. ? No definite pulmonary emboli. Results for Patient as of 12/29/2020 17:25 12/29/2020 08:19 WBC: 8.3 RBC: 3.09 (L) Hemoglobin: 10.3 (L) Hematocrit: 31.0 (L) MCV: 100.3 (H) MCH: 33.3 (H) MCHC: 33.2 RDW-CV: 15.4 RDW-SD: 57.2 (H) Platelet Count: 153 MPV: 11.5 Seg Neut Absolute: 6.9 Lymphocytes Absolute: 0.7 (L) Monocytes Absolute: 0.4 Eosinophils Absolute: 0.2 Basophil Absolute: 0.1 Neutrophils Percent: 83.5 Neutrophils Abs. (Segs and Bands): 6,900 Lymphocytes Percent: 8.8 (L) Monocytes Percent: 4.6 Eosinophils Percent: 2.1 Basophil Percent: 0.8 Glucose: 130 (H) Sodium: 138 Potassium: 4.0 Chloride: 103 CO2: 21 Anion Gap with K: 18 BUN: 17 Creatinine: 0.84 BUN/Creatinine Ratio: 20.2 Calcium: 8.8 Corrected Calcium: 9.0 Bilirubin Total: 1.1 Alkaline Phosphatase: 82 ALT - SGPT: 19 AST - SGOT: 25 Protein Total: 6.5 Albumin: 3.8 Ferritin: 168 CRP: 3.2 Lactic Acid: 1.3 eGFR African American: 75 eGFR Non-African American: 62 Protime: 17.5 (H) INR: 1.5 (H) APTT: 30.7 D-Dimer: 1.19 (H)

CDC Split Type:

Write-up: Fever, coughing, tachypnea, tachycardia

VAERS ID: 912957 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: Oklahoma
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anxiety, Chest pain, Dysphagia, Dyspnoea, Oropharyngeal discomfort, Pruritus
SMQs:, Anaphylactic reaction (narrow), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zrytec 10 mg; estradiol 1mg; atrovent; singular 10mg; ventalin Advair; Lexapro 20mg; melatonin 6 mg; vitamin D; Quercitin, Vitamin C; Vitamin B1; Zinc; ASA 81 mg; daily vitamin
Current Illness: None
Preexisting Conditions: Mast Cell Activation; Asthma
Allergies: Cats; Pork; Dilaudid; Nubain
Diagnostic Lab Data: unknown at this time
CDC Split Type:

Write-up: Received vaccine at 13 15. At 1351 c/o mild chest pain. 1353 c/o throat fullness and anxiety; 1354 bp 148/81 HR 99 O2 sat 100% c/o difficulty swallowing, SOB and itching. Rapid response called. Benadryl 50 mg IVP given at 1355. 1356 Solumedrol 125mg given IV; 1357 Epinephrine given; 1358 Pepcid 20 mg given. 1400 Normal saline 250 mg IV; 1415

transported by ambulance to local ER. In waiting area in local ER no attention. Came back to Hospital and admitted to observation

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VAERS ID: 913106 (history)
Form: Version 2.0
Age: 82.0
Sex: Female
Location: New York
Vaccinated: 2020-12-26
Onset: 2020-12-27
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood pressure increased, Breath sounds abnormal, Chills, Pneumonia, Tremor, Vomiting
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Parkinson-like events (broad), Acute central respiratory depression (narrow), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypertension (narrow), Eosinophilic pneumonia (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Amlodipine 2.5mg, aspirin 81mg, atorvastatin 10mg, bacid tablet, namenda 5mg, keppra 500mg, meclizine 12.5, mirtazapinr 15mg, tylenol 325
Current Illness:
Preexisting Conditions: Alzheimers, hyperlipidemia, thrombocytopenia, TIA, major depressive disorder
Allergies: None
Diagnostic Lab Data: BP 213/159; 92% oxygen saturation on room air
CDC Split Type:

Write-up: The morning after getting vaccine, patient had high BP 213/159, 92% on room air, shaking, chills, LCTAB with diminishment at bases, emesis. Admitted to hospital for

pneumonia.

VAERS ID: 912029 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-15
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Dizziness, Dyspnoea, Hypertension, Oxygen saturation, Paraesthesia
SMQs:, Anaphylactic reaction (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Hypertension (narrow), Cardiomyopathy (broad), Vestibular disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: No known drug allergies.
Allergies:
Diagnostic Lab Data: Test Date: 20201215; Test Name: Blood pressure; Result Unstructured Data: Test Result:170-120; Test Date: 20201215; Test Name: Oxygen saturation; Test Result: 98 %
CDC Split Type: GBPFIZER INC2020510208

Write-up: raised blood pressure, dizziness and right facial tingling, some shortness of breath; raised blood pressure, dizziness and right facial tingling, some shortness of breath; raised blood pressure, dizziness and right facial tingling, some shortness of breath; raised blood pressure, dizziness and right facial tingling, some shortness of breath; This is a spontaneous report from a contactable physician downloaded from the Agency Regulatory Authority-WEB (GB-MHRA-EYC 00235461). A 52-year-old female patient received BNT162B2 (PFIZER-

BIONTECH COVID-19 MRNA VACCINE; Lot number: EJ0553) intramuscularly on 15Dec2020 at 0.3 mL single for covid-19 immunization. The patient"s medical history was not reported. Concomitant medication included irbesartan (MANUFACTURER UNKNOWN). On 15Dec2020, the patient experienced raised blood pressure, dizziness and right facial tingling, some shortness of breath, which caused hospitalization from an unspecified date to an unspecified date. Clinical details were reported as follows: During the observation period after vaccination, patient complained of dizziness and right sided facial tingling. Some difficulty in breathing. Oxygen was started and ambulance was called and the patient was sent to hospital. The patient underwent lab tests and procedures on 15Dec2020 which included blood pressure: 170-120 and oxygen saturation: 98 %. The outcome of the events was not recovered. No follow up attempts are possible. No further information is expected.

VAERS ID: 912041 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-16
Onset: 2020-12-16
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-29

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Fatigue, Oxygen saturation, Throat tightness, Wheezing
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Asthma/bronchospasm (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Eosinophilic pneumonia (broad), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Asthma
Allergies:
Diagnostic Lab Data: Test Date: 20201216; Test Name: Oxygen saturation; Test Result: 100 %

CDC Split Type: GBPFIZER INC2020510095

Write-up: Anaphylaxis; tightness in her throat; Mild wheezing; very tired; This is a spontaneous report from two contactable pharmacists and other health care professional downloaded from the Medicines Agency (MA) regulatory authority-WEB GB-MHRA-WEBCOVID-20201216120813. Additional case identifiers: GB-MHRA-WEBCOVID-20201217103503 and GB-MHRA-WEBCOVID-20201216194631. A 25-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) lot number: EJ0553, via an unspecified route of administration on 16Dec2020 at a single dose for covid-19 immunisation. The patient''s medical history included asthma. Concomitant medication included salbutamol (metered dose inhaler), oral contraceptive nos, trazodone and beclometasone. On the 16Dec2020 the patient experienced anaphylaxis. The event was considered serious for being causing/prolonging hospitalization and being medically significant. The patient underwent lab tests and procedures which included oxygen saturation: 100% on 16Dec2020. The patient had mild wheezing post-vaccination. She was nebulised with salbutamol and her condition improved. Patient has not tested positive for COVID-19 since having the vaccine. Patient was not pregnant. She has not had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in a clinical trial. An update received which stated that the patient experienced mild wheezing with tightness in her throat. Nebulised salbutamol used in vaccine hub. Wheeze resolved and SpO2 100% on rebreathe mask. She confirmed she felt better but was very tired. Her chest was clear and patient was discharged from the vaccine hub but advised to remain with someone throughout the day and if develops further symptoms, to call 999. Patient deteriorated within 4 hours of discharge and admitted to A&E. Adrenaline and steroids administered and treated as delayed anaphylaxis. Patient admitted to hospital and feeling much better following morning. The outcome of the events Anaphylaxis and tightness in her throat was recovering; recovered for mild wheezing and unknown for very tired. No follow up attempts are possible. No further information is expected.

VAERS ID: 913218 (history)
Form: Version 2.0
Age: 25.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-27
Onset: 2020-12-27
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Dizziness, Headache, Nausea, Peripheral swelling
SMQs:, Cardiac failure (broad), Acute pancreatitis (broad), Angioedema (broad), Anticholinergic syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Haemodynamic oedema, effusions and fluid overload (narrow), Vestibular disorders (broad)

Life Threatening? No
Birth Defect? No

Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: acetaminophen 1000mg and 25mg diphenhydramine
Current Illness:
Preexisting Conditions: lupus, asthma, ulcerative colitis
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: headache, nausea, light headed, swelling of the hands Patient was given 1000mg of acetaminophen and 25mg diphenhydramine. Patient was monitored for 60 minutes.

VAERS ID: 913445 (history)
Form: Version 2.0
Age: 24.0
Sex: Female
Location: Unknown
Vaccinated: 2020-12-27
Onset: 2020-12-27
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Chest discomfort, Intensive care, Lip swelling, Pruritus, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations: Influenza vaccine = hives
Other Medications:

Current Illness:
Preexisting Conditions:
Allergies: Pt reports having hives to the influenza vaccine
Diagnostic Lab Data:
CDC Split Type:

Write-up: Pt developed anaphylaxis, was given IM Benadryl, and was sent to the ED. Pt spent 1 night in the hospital, went home, and has come back and is in the ICU. Pt had hives, itching, chest tightness, swollen lips.

VAERS ID: 913560 (history)
Form: Version 2.0
Age: 53.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	PAA 156051 / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest discomfort, Electrocardiogram normal, Myocardial necrosis marker normal
SMQs:, Anaphylactic reaction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Sertraline 100 mg twice daily
Current Illness: None
Preexisting Conditions: Lupus, RA
Allergies: PCN, Toradol, Bees, Biaxin
Diagnostic Lab Data: EKG, cardiac enzymes. All negative for cardiovascular event. 12/18/20.
CDC Split Type:

Write-up: Approximately 15 minutes after IM injection, patient developed chest tightness. Patient was taken to the Emergency Department for treatment. Was given sublingual nitroglycerin and subsequently admitted to the hospital for observation. Patient discharged

home the following morning in good condition.

VAERS ID: 913764 (history)
Form: Version 2.0
Age: 47.0
Sex: Male
Location: New Mexico
Vaccinated: 2020-12-20
Onset: 2020-12-21
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Condition aggravated, Coronary artery disease, Headache, Laboratory test, Migraine
SMQs:, Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Other ischaemic heart disease (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vit. D3, Magnesum, Acetaminophen, B12, Rosuvastatin, Trazodone, fatomadien, alloturinol, propranolol, losartan
Current Illness: migranes, diabetes hypertension
Preexisting Conditions: Lower back pain, arthritis- both knees, ankles, shoulders and thumbs, gout
Allergies: lisinopril
Diagnostic Lab Data: labs
CDC Split Type: vsafe

Write-up: About 3 am after the injection I woke up with severe chest pains and headache and went to the ER. I was admitted for 2 days and was released with a prescription of Isosorb Monoer, Metroprol, aspirin which is used as a blood thinner. The diagnosis and prognosis was severe migranes and cardioartery disease.

VAERS ID: 913807 (history)

Form: Version 2.0
Age: 49.0
Sex: Female
Location: Washington
Vaccinated: 2020-12-20
Onset: 2020-12-20
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / UNK	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Hypoaesthesia, Paraesthesia oral, Tongue dry
SMQs:, Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Dehydration (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: No
Current Illness: No
Preexisting Conditions: High blood pressure Arthritis.
Allergies: Lisinopril.
Diagnostic Lab Data:
CDC Split Type: vsafe

Write-up: Left side of my face went numb. 5 minutes later tongue was dry and tingly. Went back to place of work and went to monitoring room. Went to ER to be checked out. Got better over that day. I was admitted the same day and discharged the next day.

VAERS ID: 913866 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Minnesota
Vaccinated: 2020-12-29
Onset: 2020-12-29

Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Intensive care, Localised oedema, Pharyngeal swelling, Pruritus, Rash macular, Skin burning sensation, Throat irritation, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Prenatal Multivitamin
Current Illness: NA
Preexisting Conditions:
Allergies: Amoxicillin - Hives Iodinated Casein - Anaphylaxis Oxycodone - Hives Penicillins - Hives Roxicodone - Hives
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient administered Pfizer-BioNtech vaccine, dose #1 in series at 810AM, without notable concerns for 10 minutes. At 10 minutes post vaccination, patient developed itching and some blotching, throat becoming scratchy. No known allergies to components listed in vaccine, though does have a listed allergy to contrast dye, does not carry an epi-pen. Patient was walked to urgent/emergency care in the clinic, where she was seen immediately. Patient was given diphenhydramine 25 mg IV, pantoprazole 40 mg IV, with mild uticaria continuing. Patient notes she feels her "back is on fire". Patient was then given methylprednisolone 125 mg IV. At this time, no SOB is noted, uticaria is still present. 0930 patient reports swelling in throat and respiratory difficulties at this time with visible edema in the neck. 0.3 mg Epinephrine given at 0935, epinephrine drip and racemic epinephrine neb given. 1006 Patient on epinephrine drip 2.5 mics/hour. Patient transferred to Medical Center ICU, where she remains at the time of this report.

VAERS ID: 914139 (history)
Form: Version 2.0

Age: 46.0
Sex: Female
Location: Hawaii
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Hypoaesthesia
SMQs:, Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Facial numbness radiating down left side of neck to left arm, elbow and chest. Went to ED, admitted for observation overnight.

VAERS ID: 914280 (history)
Form: Version 2.0
Age: 63.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-17
Onset: 2020-12-26
Days after vaccination: 9
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anosmia, Blood magnesium, Brain natriuretic peptide, C-reactive protein, Chest X-ray, Chills, Cough, Diarrhoea, Dyspnoea, Electrocardiogram normal, Fibrin D dimer, Full blood count, Influenza virus test negative, Metabolic function test, Nausea, Paranasal sinus hypersecretion, Pyrexia, Red blood cell sedimentation rate, SARS-CoV-2 test positive, Troponin T
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Taste and smell disorders (narrow), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), Immune-mediated/autoimmune disorders (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: aspirin 81mg, nitro 0.4 sublingual, hydrochlorothiazide 25mg, repatha 120mg, plavix 75mg, rotonix, torsemide 5mg, zetia 10mg, zoloft 100mg, bystolic 5mg, synthroid 100mcg, telmisartan 80mg, allopurinol 100mg,
Current Illness:
Preexisting Conditions: hypothyroidism, CAD, HTN, Hyperlipidemia, GERD, gout, osa,
Allergies: Codeine, erythromycin, augmentin
Diagnostic Lab Data: COVID swab - positive EKG, - normal Flu Swabs - negative CBC, CMP, Magnesium, CRP, LFH, ESR, BNP, Tropon T, D-Dimer, Chest xray
CDC Split Type:

Write-up: Patient presented to the ER on 12/26/2020 with complaints of fever, chills and SOB for about 3 days. Stated it was a gradual onset and has been intermittent. Also reported some nausea and chronic loose stools, mild loss of smell and sinus drainage, non productive cough. Discharged to observation and then home on 12/27/2020. Patient readmitted to hospital on 12/30/2020.

VAERS ID: 914474 (history)
Form: Version 2.0
Age: 64.0
Sex: Female
Location: New Jersey

Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / UNK	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Dyspnoea, Headache, Oropharyngeal pain, Peripheral swelling, Pruritus, Rash
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (broad), Retroperitoneal fibrosis (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Acetaminophen 650 Q8, Vit C, lactobacillus 20 billions daily, metaxalone 800 mg 3 times daily
Current Illness: Asthma
Preexisting Conditions: Asthma
Allergies: Benzonatate, iodine, guaifaresin
Diagnostic Lab Data: Benadryl 50 mg then 50 IV Q8 hours, epinephrine, Pepcid,
CDC Split Type:

Write-up: headache 15 minutes after receiving vaccine, 3 to 4 hours later broke out in rash on upper extremities, ABD pain, itchy, and arm swelling, throat hurts. SOB

VAERS ID: 914479 (history)
Form: Version 2.0
Age: 41.0
Sex: Female
Location: New York
Vaccinated: 2020-12-24

Onset: 2020-12-25
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Fatigue, Hemiparesis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Noninfectious encephalitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Xyzal 5mg tab. Take 1 tab BID Amlodipine 10mg 1 tab PO QAM Kariva (Mircette) 1 tab PO QHS Mucinex (Guaifenesin) 600 mg 1tab PO BID Omega 3 Fatty acid 1cap PO QD Calcium + Vitamin D 600 mg 2 tabs PO Qhs Ocean spray BID to nostrils Nas
Current Illness: none
Preexisting Conditions: Mild to moderate Intellectual Disabilities Organic Anxiety Disorder Atypical Manic Disorder Frontal Lobe Disorder Chronic Bronchiectasis Hypothyroidism Mild sleep apnea CPAP at 6 cm Hyperlipidemia Hypertension Nasal Allergies Facial Acne Obesity , Short Stature Chronic Sinusitis Seizure Disorder
Allergies: NKA
Diagnostic Lab Data:
CDC Split Type:

Write-up: Left sided weakness, fatigue for 3 days post immunization. Patient was seen by health care provider on 12/30/2020. Provider transferred patient to Hospital ER for further evaluation.

VAERS ID: 914521 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Idaho
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Amphetamines positive, Anterograde amnesia, Antidepressant drug level, Atelectasis, Back pain, Bacterial test positive, Barbiturates positive, Blood culture, Blood lactic acid normal, Blood urine present, Chills, Cholelithiasis, Chromaturia, Coma scale abnormal, Computerised tomogram abdomen normal, Computerised tomogram head normal, Computerised tomogram pelvis, Computerised tomogram thorax normal, Condition aggravated, Cough, Culture urine, Drug screen, Dysarthria, Dysuria, Eye movement disorder, Flank pain, Gaze palsy, Glucose urine absent, Headache, Hypertonia, Hypopnoea, Incoherent, Influenza A virus test negative, Influenza B virus test, Influenza virus test negative, Intervertebral disc degeneration, Lethargy, Leukocytosis, Mental status changes, Migraine, Moaning, Myalgia, Nasal congestion, Nasal septum deviation, Nausea, Nitrite urine absent, Opiates negative, Oropharyngeal pain, Paranasal cyst, Pelvic pain, Protein urine absent, Pyrexia, Red blood cells urine negative, SARS-CoV-2 test positive, Scan with contrast normal, Seizure, Sinus disorder, Specific gravity urine normal, Unresponsive to stimuli, Urine abnormality, Urine analysis, Urine bilirubin decreased, Urine ketone body present, Urine leukocyte esterase, Urobilinogen urine, Vomiting, White blood cells urine negative, pH urine normal
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Acute pancreatitis (broad), Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhage laboratory terms (narrow), Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (narrow), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Dementia (broad), Convulsions (narrow), Parkinson-like events (narrow), Drug abuse and dependence (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (narrow), Psychosis and psychotic disorders (broad), Gallbladder related disorders (narrow), Gallstone related disorders (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Ocular motility disorders (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Proteinuria (broad), Respiratory failure (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (narrow), Hypoglycaemia (broad), Infective pneumonia (broad), Hypokalaemia (broad), Opportunistic infections (broad), COVID-19 (narrow), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none on file
Current Illness: COVID- 19 diagnosis 12/11/2020 asymptomatic
Preexisting Conditions: DIAGNOSIS 1. COVID-19 virus infection ? 2. Urinary tract infection with hematuria, site unspecified ? ? Microscopic hematuria. 3. Seizure (HCC) ? 4.

Nonintractable headache, unspecified chronicity pattern, unspecified headache type ? 5. Cough ? 6. Neutrophilic leukocytosis ? 7. Amphetamine or stimulant drug abuse (HCC) ? 8. Intracranial aneurysm ? ? Reportedly identified on MRI/MRA of brain in 2020. Unknown location, size, and characteristics. ? HX: seizures
Allergies: NKDA
Diagnostic Lab Data: Results as of 12/30/2020 14:32 12/30/2020 00:23 Color Urine: Yellow Clarity Urine: Slightly Cloudy (A) Specific Gravity: 1.010 Glucose Urine: Negative Bilirubin Urine: Negative Ketones Urine: 5 mg/dL Blood Urine: Moderate (2+) (A) pH Urine: 6.5 Protein Urine: Negative Nitrite: Negative Leukocyte Esterase Urine: Negative Urobilinogen: < 2 mg/dL WBC Urine: Negative RBC Urine: 0-2 /hpf Squamous Epithelial: Occ (0-10) /lpf Bacteria: Many ($g50) /hpf (A) 12/30/2020 00:26 Amphetamines: Positive (A) Barbiturates: Positive (A) Benzodiazepine: Negative Buprenorphine: Negative Cocaine: Negative Marijuana (THC): Negative Methadone: Negative Methamphetamines: Negative Opiates: Negative Oxycodone: Negative Phencyclidine: Negative Tricyclics Antidepressant TCA: Negative Propoxyphene: Negative 12/30/2020 00:36 Influenza B: Not Detected Influenza A: Not Detected SARS-CoV-2: Detected (A) 12/30/2020 00:58 CULTURE, BLOOD: Rpt pending 12/30/2020 01:04 CULTURE, BLOOD: Rpt pending ?Procedure: CT HEAD WITHOUT CONTRAST ?Date of Service: 12/29/2020 CT OF THE HEAD WITHOUT IV CONTRAST: ? INDICATION: Altered level of consciousness. ? COMPARISON: No prior exams are available for comparison. ? FINDINGS: No acute intracranial hemorrhage, mass effect, or obvious infarcts. ?Normal ventricular size. ?Rightward nasal septal deviation. ?Mildly prominent mucous retention cyst in the left maxillary sinus. ?Mild additional scattered paranasal sinus disease. ? The preliminary report was reviewed without significant discrepancy. ? IMPRESSION: No acute intracranial abnormalities. ?Scattered paranasal sinus disease, as discussed above. ? If clinical concern persists, MRI could be considered for further evaluation. ? ?Procedure: CT CHEST ABDOMEN PELVIS WITH CONTRAST ?Date of Service: 12/29/2020 CT OF THE CHEST, ABDOMEN, AND PELVIS WITH IV CONTRAST. ?ADDITIONAL MIP REFORMATTED IMAGES WERE REVIEWED. ? INDICATION: Leukocytosis, diffuse pain. ? COMPARISON: No prior exams are available for comparison. ? FINDINGS: CHEST: Shallow inspiration. ?Mild bibasilar dependent atelectasis. ?No focal consolidation, pleural effusion, or pneumothorax. ? Normal heart size. ?No definite pathologic lymphadenopathy. ?Benign-appearing bone island in T12 on the right. ?Mild degenerative change of the spine. ? ABDOMEN/PELVIS: Large probable noncalcified gallstone in the gallbladder neck measuring up to 1.9 x 3.0 cm with mildly prominent sludge throughout the remainder of the gallbladder lumen. ?No significant surrounding inflammatory change is identified at this time. ?Correlation with ultrasound is recommended for further evaluation. ? Liver, spleen, pancreas, adrenal glands, and kidneys appear grossly unremarkable. ?No hydronephrosis. ?No evidence for bowel obstruction or appendicitis. ?No definite pathologic lymphadenopathy. ?Degenerative changes of the spine. ? The preliminary report was reviewed with discrepancy as the large probable noncalcified stone and sludge in the gallbladder lumen was not mentioned on the preliminary interpretation. ? IMPRESSION: 1. ?Large probable noncalcified gallstone in the gallbladder neck with mildly prominent sludge, as discussed above. ?Correlation with ultrasound is recommended for further evaluation. 2. ?Additional nonacute/chronic findings, as detailed above. ?
CDC Split Type:

Write-up: Patient presents with ? Altered Mental Status ? Headache ? ? HPI Patient presents to ER by EMS ambulance after family called 911 as patient was incomprehensible with slurred speech and moaning on the phone this evening. On arrival of EMS patient was asleep in bed and reportedly unresponsive other that to localize to pain. EMS transferred patient to ER. On arrival to ER patient had GCS 7. Reportedly patient is locum nurse who works in a Nursing Home and patient reportedly received COVID vaccination 2 days ago and that night reportedly began complaining to family on the phone of headache, nasal congestion, sore throat, cough, fever, chills, nausea, emesis, myalgias, and lethargy. Per the medical record patient has

history of seizures, migraines, and sciatica. No other information is known on patient arrival to ER. 1. Peripheral IV right dorsal hand placed by EMS in route to ER. 2. On arrival to ER GCS 7 (E1M5V1) and roving eye movements with episodic lateral conjugate and at times disconjugate gaze concerning for seizure activity. Arms and legs with moderately increased tone but no clonic movements and patient able to localize bilaterally. 3. Ativan 1 mg IVP for seizure, then further 2 mg IVP for persistent seizure. 4. Fosphenytoin 1,000 mg IVPB in ER for loading dose of antiseizure medication. 5. Patient had significant improvement following completion of Ativan 3 mg IVP and GCS improved to 14 (E3M6V5) from GCS of 7 (E1M5V1). 6. Patient able to communicate after improvement as above and reports she has had headache or migraine for past several days as well as dysuria with bilateral CVA pain and has significant pain on percussion of bilateral CVA and moderate pain on palpation of bilateral flanks. No nuchal rigidity or pain with ROM of neck. Additionally, she complains of severe headache and diffuse pain of back and abdomen/pelvis. She reports a history of seizures in the past and reports she had one last month and was treated at a hospital in her home state. She denies antiseizure medications. Additionally, patient reports nonproductive cough, sore throat, nasal congestion, fever, chills, myalgias, lethargy, nausea, and episodic emesis over the past 2 days. She has anterograde amnesia following seizure and does not recall events. 7. Normal saline 1,000 mL IV bolus, then 100 mL/hour in ER. 8. CT of head with and without contrast performed and negative for intracranial hemorrhage, lesions, stroke, or other acute pathology. 9. CT of chest/abdomen/pelvis with IV contrast shows no acute pathology or notable abnormalities. 10. Lactic acid drawn and normal. 11. UA and urine microscopy collected by straight catheterization and culture collected and pending. 12. Blood cultures x 2 collected and pending. 12. Rocephin 2 mg IVPB in ER after blood cultures collected. 13. Vancomycin 20 mg/kg (1,500 mg) IVPB following Rocephin. 14. Dexamethasone 10 mg IVP in ER. 15. Duoneb nebulizer in ER. 16. Called to discuss with patient''s daughter and the family''s preferred contact who is 23 years old. She reports patient had COVID vaccination 2 days ago and beginning that night patient has complained of headache, fever, chills, nonproductive cough, sore throat, nasal congestion, myalgias, and lethargy. Additionally, she reports patient has history of seizures on at least 1 occasion in the past a few months ago but is not on antiseizure medications. She reports patient had MRI and MRA of her brain at that time and reportedly and intracranial aneurysm was identified at that time. 17. Called to request transfer to another facility and spoke with hospitalist who states there is no neurologist there and suggests transfer to larger tertiary care facility with neurology. 18. Called to request transfer and accepted by ER provider. 19. Transfer by ALS ground ambulance with telemetry, pulse oximetry, O2 to keep $g 02%, vitals every 30 minutes, normal saline at 100 mL/hour, Rocephin 2 grams IVPB, vancomycin 20 mg/kg (1,500 mg) IVPB in ER. ? DISPOSITION Patient Stabilized and Transferred Data Unavailable Wed Dec 30, 2020 2:12 AM CST

VAERS ID: 914730 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: California
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site /

		Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram coronary artery normal, Dizziness, Electrocardiogram T wave inversion, Full blood count normal, Hyperhidrosis, Metabolic function test normal, Myocardial necrosis marker increased, Nausea, Presyncope, Troponin normal
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (broad), Anticholinergic syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Other ischaemic heart disease (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), Hypokalaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Bupropion HCl XL 300mg tablet QDay Bupropion HCl SR 100mg tablet QDay Escitalopram 20mg tablet QDay Buspirone HCL 7.5mg tablet TID
Current Illness: None
Preexisting Conditions: Scoliosis Depression
Allergies: None
Diagnostic Lab Data: EKG 12/17/2020, inverted T waves CBC, BMP, within normal limits Cardiac enzymes x 3, troponin was zero Coronary CT with contrast, WNL
CDC Split Type:

Write-up: Near syncopal episode approximately 2.5 hours after vaccination. Sudden onset of dizziness, nausea, and diaphoresis. Was admitted to ED and observed overnight. Full cardiac work up was done and shown to be within normal limits. I have no pre-existing conditions and considered to be a healthy adult.

VAERS ID: 914798 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: California
Vaccinated: 2020-12-18
Onset: 2020-12-20
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site /

		Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Bundle branch block right, Cardiac stress test normal, Chest X-ray, Chest pain, Depressed level of consciousness, Electrocardiogram abnormal, Full blood count, Heart rate decreased, Hyperhidrosis, Metabolic function test, Nausea, Pallor, Sinus bradycardia, Transcutaneous pacing, Troponin increased
SMQs:, Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Disorders of sinus node function (narrow), Conduction defects (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Amlodipine 10 mg daily
Current Illness: none
Preexisting Conditions: Hypertension
Allergies: No known allergies
Diagnostic Lab Data: EKGs, Chest X-rays, Serial Toponin Levels, CBC, CMP, Cariac Stress test
CDC Split Type:

Write-up: On Dec. 20, 2020 around 11:30 PM, 2 days after patient received her COVID-19 vaccination, she was found on the bathroom floor , obtunded, very pale, diaphoretic, nauseous, and complaining of severe chest pain. Paramedics was called and patient was transported to the nearest emergency room. According to paramedics, on the way to the ER while patient was in the ambulance,she was noted with a sudden drop in heart rate about 19 beats/minute and have to be given Atropine IV Push, oxygen and was connected to transcutaneous pacing which improves her heart rate. In the ER patient continued to have chest pain and she was given Morphine, Oxygen, Nitroglycerine and Aspirin. IM had an EKG which showed Sinus Bradycardia with a Right Bundle Branch Block. She had serial ekgs, a chest x-ray, laboratory testing which included Troponin. Her first Troponin level came back elevated prompting her hospital admission to Telemetry. Her next 2 Troponin level improved and return to normal range and her chest pain has resolved.. She underwent a Stress Test which came back negative. Patient was admitted for a total of 20 hours in the Telemetry unit with Cardiology consultation before being discharged home last . She was re-evaluated by the cardiologist yesterday which diagnosed her a chest pain of unknown origin.

VAERS ID: 914835 (history)
Form: Version 2.0
Age: 51.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-26
Onset: 2020-12-28
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0142 / 1	- / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Dyspnoea, Hypoxia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Went to Emergency room on 12/28/20 because he was short of breath. Tested positive for COIVD_19 and was admitted with hypoxia.

VAERS ID: 915119 (history)
Form: Version 2.0
Age: 58.0
Sex: Male
Location: Wyoming
Vaccinated: 2020-12-24

Onset: 2020-12-29
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: C-reactive protein increased, COVID-19 pneumonia, Chest X-ray abnormal, Computerised tomogram thorax abnormal, Dyspnoea, Fibrin D dimer increased, Hypoxia, SARS-CoV-2 test negative
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Haemorrhage laboratory terms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: albuterol, Zithromax, Advair
Current Illness: diagnosed with covid19 infection on 12/14/2020, had ulnar nerve and median nerve releases in November, 2020
Preexisting Conditions: tobacco abuse.
Allergies: no known allergies
Diagnostic Lab Data: CT of chest and CXR supported covid19 pneumonia diagnosis with marked elevation in D-dimer, CRP. interestingly SARS-CoV2 rapid antigen was negative on 12/29/2020.
CDC Split Type:

Write-up: The patient had COVID19 infection diagnosed 12/14/2020, and he stated 5 to 10 days after this, he developed shortness of breath. Had vaccine on 12/24/2020. Hypoxic and short of breath with COVID19 pneumonia on 12/29/2020. I do not know if this is an adverse effect or temporally related or if the vaccine activated prior infection.

VAERS ID: 913148 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 0000-00-00
Onset: 2020-12-10

Submitted: 0000-00-00
Entered: 2020-12-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Cold sweat, Dizziness, Headache, Myalgia, Nausea, Pain, Pallor, Peripheral coldness, Somnolence, Syncope, Tremor
SMQs:, Torsade de pointes/QT prolongation (broad), Rhabdomyolysis/myopathy (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Parkinson-like events (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19 (had COVID-19 and was very ill with it.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2020514800

Write-up: Cold sweat; Dizziness; Nausea; Pallor; Peripheral coldness; Somnolence; Tremor; Fainted after having jab and now having all the side effects, myalgia,severe body aches , headaches; Fainted after having jab and now having all the side effects, myalgia,severe body aches , headaches; Fainted after having jab and now having all the side effects, myalgia,severe body aches , headaches; Fainted after having jab and now having all the side effects, myalgia,severe body aches , headaches; This is a spontaneous report from a contactable consumer, downloaded from the Regulatory Authority GB-MHRA-EYC 00235088, Safety Report Unique Identifier GB-MHRA-ADR 24542268, EU-EC-10007183035. A 37-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date at a single dose for COVID-19 vaccination. Medical history included COVID-19 and was very ill with it. Concomitant medication included ibuprofen, taken from an unknown date for unknown indication. The patient experienced fainted after having jab and now having all the side effects, myalgia, severe body aches, and headaches on 10Dec2020. On an unknown date, the patient also experienced cold sweat, dizziness, headache, nausea, pallor, peripheral coldness, somnolence, and tremor and these events were reported to have caused hospitalization and

other medically important condition. The outcome of fainted after having jab and now having all the side effects, myalgia, severe body aches, and headaches was not recovered, dizziness was recovering, while the outcome of the other events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 915161 (history)
Form: Version 2.0
Age: 34.0
Sex: Female
Location: Oregon
Vaccinated: 2020-12-18
Onset: 2020-12-19
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Breast pain, Breast tenderness, Headache, Lymph node palpable, Lymphadenopathy, Musculoskeletal stiffness, Pain, Pain in extremity, Tenderness
SMQs:, Dystonia (broad), Parkinson-like events (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Lipodystrophy (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020506596

Write-up: my left arm and breast were tender with some mild body aches and low grade headache/Arm soreness; my left arm and breast were tender with some mild body aches and low grade headache/ left breast was increasingly painful with swollen palpable lymph nodes; my left arm and breast were tender with some mild body aches and low grade headache/ left breast was increasingly painful with swollen palpable lymph nodes; my left arm and breast

were tender with some mild body aches and low grade headache/ left breast was increasingly painful with swollen palpable lymph nodes; my left arm and breast were tender with some mild body aches and low grade headache; body soreness and stiffness; This is a spontaneous report from a contactable healthcare professional. A 34-year-old female patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EH9899), via an unspecified route of administration on 18Dec2020 at single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient stated, "the day after the injection, my left arm and breast were tender with some mild body aches and low grade headache. Sunday Arm soreness went away but left breast was increasingly painful with swollen palpable lymph nodes, with body soreness and stiffness. Monday left breast is still remarkably tender with palpable lymph nodes" on 19Dec2020 (reported as Seriousness criteria-Caused/prolonged hospitalization: Yes, on an unspecified date in Dec2020). The outcome of the event was not recovered.; Sender"s Comments: The reported information is limited. Based on the close temporal relationship and the description of the events, there is a reasonable possibility that the events are related to BNT162 vaccine. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 915478 (history)
Form: Version 2.0
Age: 55.0
Sex: Male
Location: Georgia
Vaccinated: 2020-12-30
Onset: 2020-12-31
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dystonic tremor, Erythema, Headache, Tachycardia, Tachypnoea
SMQs:, Anaphylactic reaction (narrow), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dystonia (narrow), Eosinophilic pneumonia (broad), Hypersensitivity (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: Crestor, Allopurinol
Current Illness: N/A
Preexisting Conditions: Recovered from COVID 19. Positive on 9/11/2020
Allergies: Iodine
Diagnostic Lab Data:
CDC Split Type:

Write-up: erythema, tachycardia, tachypnea, headache, uncontrolled dystonic shaking

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VAERS ID: 915480 (history)
Form: Version 2.0
Age: 75.0
Sex: Female
Location: Minnesota
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	- / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Chest X-ray normal, Chills, Injection site erythema, Injection site induration, Laboratory test normal, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vitamin D3 4,000 units daily, famotidine 20 mg daily, ferrous sulfate 325 mg daily, gabapentin 400 mg BID, levothyroxine 100 mcg daily, melatonin 9 mg at hs, metoprolol tartrate 12.5 mg BID, nortriptyline 10 mg at HS, Norvasc 10 mg daily, S
Current Illness: Had a uti 12/8/2020 finished abx on 12/10/2020; prior to admitting had COVID with multiple complications spent along time in the hospital recovering.
Preexisting Conditions: Type 2 diabetes, morbid obesity, hypertension, sleep apnea, anemia, GERD, chronic pain
Allergies: Benzalkonium, lisinopril, Percocet, Adhesive Tape
Diagnostic Lab Data: Sent to ER admitted to hospital under observation. Per Hospital notes

"There does not appear to be a cellulitis or abscess at this time. Labs and chest x-ray in the ED did not reveal any origin to where her fevers may be coming from".
CDC Split Type:

Write-up: Developed sudden onset of shaking chills and fevers as high as 103.0. She has developed a small circular 5 x 5 area of erythema and firmness at the injection site of her left upper arm.

VAERS ID: 915613 (history)
Form: Version 2.0
Age: 33.0
Sex: Male
Location: Illinois
Vaccinated: 2020-12-17
Onset: 2020-12-19
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ685 / 1	LA / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Angioedema, Blood glucose increased, Hyperglycaemia
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Hyperglycaemia/new onset diabetes mellitus (narrow), Oropharyngeal allergic conditions (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: angiodema. hospitalized overnight. During the hospitalization pt was started on an epi gtt and given MTP 125 mg. He had subsequent hyperglycemia, and increased his rate on his insulin pump to 2 U/hr (from 0.83 U/hr). Pt then decreased his rate to 1.4 U/hr while on the epi gtt, and then to 1.1 U/hr when off of the epi gtt. He also strengthened his carb ratio from

1:12 to 1:10 last night.Pt reports he had postprandial hyperglycemia overnight after his meal, but then BG corrected overnight. Pt reports fasting this morning is in the 110s. This morning"s breakfast on 1:10 CR with well controlled BG.

VAERS ID: 915690 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Consciousness fluctuating, Discomfort, Exposure during pregnancy, Fatigue, Headache, Nausea, Palpitations
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Maxalt 10mg Elavil 10mg
Current Illness: Anxiety
Preexisting Conditions:
Allergies: NKDA
Diagnostic Lab Data:
CDC Split Type:

Write-up: Due Date unknown/6-7 wks pregnant -nausea -Body felt heavy -headache -in & out consciousness -Fatigue, severe -Heart Racing

VAERS ID: 915813 (history)
Form: Version 2.0
Age: 61.0
Sex: Male
Location: Texas
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dysphagia, Full blood count normal, Metabolic function test normal, Paraesthesia oral, SARS-CoV-2 test negative, Swollen tongue
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Anticholinergic syndrome (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Hydralazine 50mg BID last taken 3 days prior to vaccination Carvedilol 12.5mg BID last taken 3 days prior to vaccination Aspirin 81mg taken 12/30 AREDS multivitamin taken 12/30
Current Illness:
Preexisting Conditions: Macular degeneration HTN
Allergies: Previous experienced three episodes of angioedema. One episode was thought to be from losartan. Patient experienced 2 additional episodes while off losartan. Previously diagnosed with seasonal allergies. Patient stated all three episodes he had eaten some sort of salad/vegetable and thinks this could be related to environmental allergies.
Diagnostic Lab Data: CMP and CBC were unremarkable 12/30 COVID-19 PCR was negative 12/30
CDC Split Type:

Write-up: Patient stated he stopped his blood pressure medications 3 days prior to vaccination due to a previous reaction to losartan, a medication he was no longer taking. Patient took aspirin and a MVI on day of vaccination and drank lemon water. Patient developed tingling sensation in his mouth after eating dinner around 18:00. Patient stated he ate tacos with apple cider and noticed tingling after dinner. Patient stated he took two benadryl with no relief. His tongue continued to swell and he took two additional benadryl at 22:00. Once he developed difficulty swallowing he went to the emergency department. Patient presented to the ED with tongue swelling and difficulty swallowing. At 23:57 he was adminsitered 0.3mg of epinephrine IM, diphenhydramine 25mg IV, famotidine 40mg IV, dexamethasone 10mg IV at 0114, methylprednisolone 60mg q6hrs started at 0417, diphenhydramine 25mg q6hrs IV started at 0416, albuterol 2.5mg via neb q6hrs started at 0710

VAERS ID: 915928 (history)
Form: Version 2.0
Age: 38.0
Sex: Female
Location: Missouri
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Confusional state, Dizziness, Dysphagia, Dyspnoea, Endotracheal intubation, Immediate post-injection reaction, Palpitations, Paraesthesia oral, Pruritus, Rash, Rash erythematous, Tremor, Vision blurred, Wheezing
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (narrow), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Glaucoma (broad), Cardiomyopathy (broad), Lens disorders (broad), Eosinophilic pneumonia (broad), Retinal disorders (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? Yes
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Percocet, reglan, and emgality migrain injection
Diagnostic Lab Data:
CDC Split Type:

Write-up: Started feeling a reaction immediately after the vaccine, felt blurred vision, dizziness, racing heartbeat, chest rash and face, itching all over, difficulty swallowing, tongue tingling and wheezing. Sent to ED. EPI and Benadryl. 1800 Went to see her in the ED, room 33. She has red rash to neck, shaky hands itching to neck and chest. ED Dr to discharge, she stated husband to pick her up and she will follow up with OH tomorrow.
----------RN
ED gave her Epinephrine 0.3 mg, Methylprednisolone 125mg, Diphenhydramine HCL 50 mg, Zofran 4mg, Lorazepam 1 mg, Hydroxyzine HCL 50 mg Sumatriptan 6mg , Discharge from ED at 1902

- RN 12/29/2020 1715 called to check on patient. left voicemail for her to call OH. ???????..? 12/29/2020 1838 left voicemail for patient to call OH. ??????????????????????. 12/30/20 2030 spoke with her. Tuesday 12/29 3pm-4pm dizziness, confusion, sob. Wheezing. Ambulance called. Hospital admitted. Intubated for less than 24 hours. Breathing treatments, epi drip. Now just on steroids and walking around and feeling better. Still admitted at hospital. Hoping discharged tomorrow. ----------RN

VAERS ID: 915956 (history)
Form: Version 2.0
Age: 47.0
Sex: Male
Location: New Jersey
Vaccinated: 2020-12-24
Onset: 2020-12-25
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / UNK	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: COVID-19, SARS-CoV-2 test positive
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: none
Preexisting Conditions:
Allergies: nkda
Diagnostic Lab Data: COVID PCR + 12/27/2020
CDC Split Type:

Write-up: Patient received dose 1 of COVID vaccine 12/24. He developed symptoms consistent with COVID infection on 12/25. He was seen in the emergency room at Hospital on 12/27,was diagnosed as COVID positive, and was discharged to home. He returned to the emergency room on 12/29 and was admitted to the hospital for treatment related to COVID infection. He is currently admitted to Hospital.

VAERS ID: 916042 (history)
Form: Version 2.0
Age:
Sex: Female
Location: California
Vaccinated: 2020-12-28
Onset: 2020-12-31
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Blood test, Dizziness, Feeling abnormal, Hypoaesthesia, Paraesthesia
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Dementia (broad), Guillain-Barre syndrome (broad), Vestibular disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations: Flu vaccine in Sept 2020 - Sore arm
Other Medications: ibuprofen, tylenol
Current Illness: no COVID- August 2020

Preexisting Conditions: no
Allergies: sulfa
Diagnostic Lab Data: Went to ED and don"t remember which one They did run blood work.
CDC Split Type:

Write-up: 10 minutes after vaccine numbness and tingling in left foot, hand and left side of face. Spacing out feeling. Feel like i"m going to pass out.

VAERS ID: 916065 (history)
Form: Version 2.0
Age: 37.0
Sex: Female
Location: Washington
Vaccinated: 2020-12-27
Onset: 2020-12-28
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / SYR

Administered by: Other **Purchased by:** ?
Symptoms: Exposure during pregnancy, Foetal monitoring, Hydrops foetalis, Premature labour
SMQs:, Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow), Foetal disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zofran 2weeks prior, Multivatamins
Current Illness: none
Preexisting Conditions:
Allergies: Sulfa, Vicodin, Neosporin
Diagnostic Lab Data:
CDC Split Type:

Write-up: EDD - 4/1/2021 - Contractions at 26 w 3 days sent to L&D to be monitored on 12/28/20. Covid-19 Sars Vaccine given 1st dose 12/27/20. Patient was diagnosed with Fetal Hydrops. Patient Hospitalized 12/28/20 - current MFM consulting in hospital & outpatient

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VAERS ID: 916247 (history)

Form: Version 2.0
Age: 32.0
Sex: Female
Location: Arkansas
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / UNK	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood pressure increased, Body temperature increased, Breast oedema, Injection site erythema, Injection site oedema, Localised oedema, Oedema peripheral
SMQs:, Cardiac failure (broad), Angioedema (broad), Neuroleptic malignant syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypertension (narrow), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Redness about 1 1/2 inches around injection site, edema to right arm, r side of neck, r breast, r shoulder, temp 99.3, increased blood pressure. Patient went to primary care physician. She was then sent to the ER. She was Covid (+) in July 2020

VAERS ID: 916473 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-31

Onset: 2020-12-31
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Cardiac assistance device user, Dyspnoea, Headache, Loss of consciousness, Seizure, Unresponsive to stimuli
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Convulsions (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: COPD, depression, pace-maker, diabetes, obesity, osteoarthritis, plantar fasciitis, localized edema, moderate persistent asthma, sleep apnea, and ovarian cyst.
Allergies: Penicillin and Blueberries
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient complained of a headache, then patient was losing consciousness and gasp for air. EpiPen was utilized due to being unresponsive then began to seizing. Patient started to flatline and an AED was used while a paramedic came. Patient was put on her side then facility gave something for seizures. Lastly, ,ambulance took her to the hospital.

VAERS ID: 916497 (history)
Form: Version 2.0
Age: 65.0
Sex: Female
Location: Massachusetts
Vaccinated: 2020-12-27

Onset: 2020-12-28
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L2OA / 1	- / SYR

Administered by: Private **Purchased by:** ?
Symptoms: C-reactive protein increased, Catheterisation cardiac normal, Chest discomfort, Chills, Coronary artery disease, Echocardiogram abnormal, Ejection fraction decreased, Haematocrit normal, Haemoglobin normal, Hypokinesia, Left ventricular end-diastolic pressure increased, Myalgia, Nausea, Red blood cell sedimentation rate increased, Stress cardiomyopathy, Troponin increased, Ventricular hypokinesia, White blood cell count normal
SMQs:, Rhabdomyolysis/myopathy (broad), Cardiac failure (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (narrow), Eosinophilic pneumonia (broad), Other ischaemic heart disease (narrow), Hypotonic-hyporesponsive episode (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vitamin D3, omega-3, Vitamin A/C, psyllium
Current Illness: none
Preexisting Conditions: History of Renal Cell Carcinoma s/p nephrectomy, Hyperlipidemia, osteopenia, sinus bradycardia, cervical radiculopathy
Allergies: None
Diagnostic Lab Data: TTE 12/30: EF 35%. Regional variation with normal contraction at the base and hypokinesis of the mid and distal segments with perhaps some sparing of the apex. Pattern is consistent with Takotsubo cardiomyopathy. Cardiac Catheterization 12/30: Mild Coronary Disease in LAD. Otherwise minimal CAD. Mildly elevated left sided filling pressure. ESR: 24 CRP <5 WBC 6.46, H/H 13.7/39.7
CDC Split Type:

Write-up: Patient started having myalgia, chills, nausea on the next day of the vaccination. on 2nd day (12/29) patient had chest pressure which made her present to Hospital ED. She had troponin elevation to 1.14. Cardiac Catheterization was done which was negative. On Trans Thoracic Echocardiogram, patient was found to have hypokinesis of the mid and distal segment with some sparing of apex proving Takotsubo (stress induced) cardiomyopathy. Patient did not have any underlying emotional or physical stress going on in her life or family. Till now extensive infectious as well as inflammatory work up is done to rule out any secondary causes of cardiomyopathy which till date have remained negative. As a diagnosis of exclusion, her presentation seems to be COVID-19 vaccine induced Takotsubo Cardiomyopathy

VAERS ID: 916508 (history)
Form: Version 2.0
Age: 81.0
Sex: Male
Location: Arizona
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2020-12-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Acute kidney injury, Blood creatinine increased, Blood glucose increased, Blood potassium decreased, C-reactive protein increased, COVID-19 pneumonia, Chest X-ray abnormal, Cough, Decreased appetite, Hyperglycaemia, Hypokalaemia, Hypoxia, Intensive care, Malaise, Pneumonia, SARS-CoV-2 test positive, Troponin increased
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Hyperglycaemia/new onset diabetes mellitus (narrow), Myocardial infarction (narrow), Retroperitoneal fibrosis (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Dehydration (broad), Hypokalaemia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: escitalopram 10mg QD lisinopril 10mg QD trazodone 100mg QHS atorvastatin 20mg HS bupropion XL 150mg QD empagliflozin 10mg QD insulin 70/30 kwikpen 22units QAM, 20units QPM
Current Illness: COVID + pneumonia (this was undiagnosed at the time of vaccination b/c he had no symptoms upon vaccination prescreening)
Preexisting Conditions: Hypertension Benign prostatic hyperplasia Depression Uncontrolled diabetes mellitus Hyperlipidemia Latent tuberculosis
Allergies: Benzoin tincture
Diagnostic Lab Data: Pt is an 81 year old male with pmedhx of uncontrolled DM2, HTN,

depression, BPH, HLD, insomnia, hx of EtOH w/d seizure. Pt states he called EMS because he had not had an appetite for 3 days and had not been eating much. EMS noted pt''s SpO2 was 62% on RA, increased to 90s after several min on 15L NRB. Pt denied shortness of breath at any point, denied sweats/chills, HA, vomiting/diarrhea. He did states he had a cough in addition to loss of appetite. CXR with finding of BL PNA, Abbott COVID + in ER. CRP 31.4. Pt was also noted to be hyperglycemic with glucose 408, trop elevated 0.12 but without chest pain, AKI creat 1.4 with baseline 0.9, and hypokalemic with K 3.0. Pt admitted to ICU for COVID related hypoxia. Notably, pt was vaccinated for COVID 19 with 1st Moderna vaccine today in am. Pt states he told screeners he felt well because he did not have shortness of breath or body aches. As of 12/31/2020, patient is still hospitalized in ICU.
CDC Split Type:

Write-up: Pt received 1st dose of Moderna vaccine in am at COVID vaccination clinic. On presentation to clinic he stated he was feeling well. Pt was brought to ER in pm with hypoxic, requiring 15L supplemental O2. Per pt''s family, pt was not feeling well for the last couple of days but didn''t think it was related to COVID. Abbott COVID + in ER. Possible vaccine reaction, though seems unlikely.

VAERS ID: 916710 (history)
Form: Version 2.0
Age: 23.0
Sex: Female
Location: Missouri
Vaccinated: 2020-12-29
Onset: 2021-01-01
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Appendicitis, Band neutrophil percentage increased, Surgery, White blood cell count increased
SMQs:, Neuroleptic malignant syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Synthroid
Current Illness:
Preexisting Conditions: Hypothyroidism

Allergies: NKDA
Diagnostic Lab Data: WBCs 13k, 4% Bands, CT with acute appendicitis- taken to surgery
CDC Split Type:

Write-up: Acute appendicitis, onset morning of 1/1/2021 (Reporting this because Pfizer covid vaccine had 3-4x higher risk of appendicitis, although data not reported for Moderna covid vaccine)

VAERS ID: 916809 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Washington
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blepharospasm, Chest discomfort, Computerised tomogram, Dizziness, Dyspnoea, Limb discomfort, Paraesthesia, Pyrexia, Tremor
SMQs:, Anaphylactic reaction (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dystonia (broad), Parkinson-like events (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Periorbital and eyelid disorders (narrow), Ocular motility disorders (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 9 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: NA
Current Illness: None
Preexisting Conditions: NA
Allergies: None
Diagnostic Lab Data: 12/28/2020 CT SCAN 12/28/2020 PULMONOLOGY CONSULT

CDC Split Type:

Write-up: CAREGIVER RECEIVED FIRST VACCINE DOSE AND SOON AFTER BEGAN TO FEEL DIZZY AND HER EYES BEGAN TO TWITCH, FOLLOWED BY UNCONTROLLED SHAKING, WITH HIGH FEVER FOLLOWED BY SEVERE SHORTNESS OF BREATH AND GASPING, WITH TIGHTNESS AROUND THE CHEST. TRANSPORTED TO EMERGENCY DEPARTMENT. HAD MULTIPLE EPISODES OF ITEMS LISTED ABOVE WHILE IN ED, INCLUDING HEAVINESS IN HER LEGS AND TINGLING IN ARMS. SHE WAS DISCHARGED FROM ED AT 10:30PM BUT WAS READMITTED ON 12/24 TO ED FOLLOWING SIMILAR ISSUES. TO DATE SHE HAS HAD 5 RAPID RESPONSES IN HOSPITAL DUE TO REPEAT OF SIGNS/SYMPTOMS. CT AND PULMONOLOGY CONSULT SCHEDULED FOR 12/28/2020.

VAERS ID: 916836 (history)
Form: Version 2.0
Age: 55.0
Sex: Male
Location: Maryland
Vaccinated: 2020-12-24
Onset: 2020-12-30
Days after vaccination: 6
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest X-ray abnormal, Chest discomfort, Computerised tomogram thorax abnormal, Coronary artery disease, Diarrhoea, Discomfort, Echocardiogram normal, Ejection fraction normal, Electrocardiogram Q wave abnormal, Electrocardiogram abnormal, Electrocardiogram change, Epistaxis, Fatigue, Fibrin D dimer normal, Flushing, Full blood count normal, Glucose tolerance impaired, Glycosylated haemoglobin increased, Injection site pain, Lipase normal, Liver function test normal, Lung opacity, Metabolic function test normal, SARS-CoV-2 test negative, SARS-CoV-2 test positive, Sinus tachycardia, Troponin normal, Urine analysis normal
SMQs:, Anaphylactic reaction (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Hyperglycaemia/new onset diabetes mellitus (narrow), Interstitial lung disease (narrow), Myocardial infarction (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Pseudomembranous colitis (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Other ischaemic heart disease (narrow), Lipodystrophy (broad), Hypersensitivity (broad), Noninfectious diarrhoea (narrow), Infective pneumonia (broad), Dehydration (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: No current medications
Current Illness: Pt reported frequent urination for several months
Preexisting Conditions: None
Allergies: No known allergies
Diagnostic Lab Data: SARS-CoV-2 RNA - Detected A 12/30/2020 @1825
CDC Split Type:

Write-up: Patient is a 55 year old male with no past medical history who presents with complaint of sudden onset of left-sided nonradiating chest discomfort of sudden onset approximately 1 hour prior to presentation while doing administrative work at rest. He describes the pain as a dull heaviness sensation and approximate 3/10 pain severity. Chest discomfort was associated with a feeling of flushing that was quite transient but chest discomfort was persistent. Patient immediately presented to the ED for further evaluation. He denies experiencing any chest pain upon waking up this morning. Does note that he did have a transient episode of epistaxis on his way to work for which he had to pull over and apply pressure to his nose but this subsequently subsided and he attributed this to dry air as he has experienced epistaxis in the past but with less severity previously. In the ER, vital signs noted for BP 133/76, pulse ranging 91-114, respiratory rate 16-20, 96% on room air. Initial laboratory parameters were completely normal including normal CBC, CMP, LFT, lipase, UA, and normal D-dimer. Initial troponin was negative x1. EKG with sinus tachycardia, heart rate of 115. Noted Q waves inferiorly. No acute ST or T wave changes appreciated. Chest x-ray with mild increased density in the left lower lobe. Given this, patient was tested for rapid Covid which was negative but PCR was positive for COVID. CT of the chest noted for focal subsegmental groundglass infiltrate at the superior segment of the LLL, likely infectious versus inflammatory. Also noted small nonspecific groundglass attenuation with focal septal thickening at the right upper lobe which could be infectious or inflammatory, bibasilar atelectasis. Patient was treated with aspirin 324 mg in the ED. Of note, patient actually just received the COVID-19 vaccination on 12/24/20. He denies any shortness of breath, no cough, denies any nausea or vomiting, denies any change in taste or smell nor change in appetite. Does note 1 single episode of loose stool but otherwise denies any diarrhea. Does report that he had approximate 48 to 72-hour period of fatigue and soreness at the site of the left deltoid injection following the vaccination but otherwise no further symptoms. It is also noted that he does have a positive family history of coronary artery disease as his dad had an MI at the age of 49. Patient has never undergone a cardiac catheterization in the past but does report having a negative stress test at the age of 42. He is being admitted under the hospitalist service for further management Patient was initially admitted under observation for chest pain obs. However patient''s Covid test came back positive and patient also had dynamic EKG changes concerning for possible unstable angina. Patient was treated with aspirin Plavix full-strength Lovenox along with beta-blocker and a cardiology consult. Serial troponins were negative. Echocardiogram revealed normal EF of 55 to 60% with no hemodynamically significant valvular disease. Cardiology felt that patient likely has underlying coronary artery disease have recommended discharge home with aspirin and Plavix with outpatient stress testing given his positive Covid testing. At the time of discharge patient denied any chest pain or shortness of breath. Patient was borderline diabetic with a hemoglobin A1c of 6.1. Patient was discharged home with Metformin along with glucometer, glucose strip, lancets. Given patient''s tachycardia patient''s Metformin 25 mg twice daily was changed to Toprol 25 mg daily. (Please note clarification in comparison to discharge home med list. Toprol XL 25 mg daily was called to pharmacy in place of the metoprolol.)

VAERS ID: 916890 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-01
Onset: 2020-12-22
Days after vaccination: 21
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Chest X-ray, Dyspnoea, Laboratory test, SARS-CoV-2 test, Throat tightness, Urticaria, Wheezing
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Asthma/bronchospasm (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zoloft, vitamin d, vitamin b, zinc, Wellbutrin, singular
Current Illness: None
Preexisting Conditions: None until vaccine
Allergies: None
Diagnostic Lab Data: COVID TEST, LABS, Chest-X-Ray
CDC Split Type:

Write-up: HIVES, SOB, THROAT CLOSING UP, WHEEZING

VAERS ID: 917026 (history)
Form: Version 2.0
Age: 82.0
Sex: Male

Location: Louisiana
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J208 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Cerebrovascular accident, Drooling, Dysphagia, Facial paralysis, Gastrostomy, Mobility decreased, Musculoskeletal disorder
SMQs:, Rhabdomyolysis/myopathy (broad), Anticholinergic syndrome (broad), Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Dyskinesia (broad), Dystonia (broad), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hearing impairment (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Codeine Peanuts
Diagnostic Lab Data:
CDC Split Type:

Write-up: 12/28/2020, Pharmacy staff administered Moderna COVID Vaccine. 12/29/2020, he had not eaten breakfast or lunch but did consume fluids and take his medications. BP =150/70, Temp. = 101.6, Pulse= 102, Respirations= 18 and Oxygen saturation= 97%. Tylenol 650 mg given. It was difficult for him to swallow. Also had no use of right upper extremity and unable to move lower extremity, mouth was drooping and was drooling. Physician in attendance and ordered to send to ER. 1/1/2021, received information from nurse at hospital that patient received a Peg Tube this afternoon and Clinical indication of a stroke.

VAERS ID: 917042 (history)
Form: Version 2.0
Age: 43.0

Sex: Male
Location: Michigan
Vaccinated: 2020-12-27
Onset: 2020-12-28
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram, Electrocardiogram, Electroencephalogram, Loss of consciousness, Magnetic resonance imaging, Nausea, SARS-CoV-2 test negative, Seizure, Vomiting
SMQs:, Torsade de pointes/QT prolongation (broad), Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: EKG 12/28, CT 12/28, MRI 12/29, EEG 12/28-12/29, covid-19 test 12/28 (negative).
CDC Split Type:

Write-up: I received the vaccine at 6:30pm on 12/27. I worked atb7pm and took lunch at 2:00am. After lunch, I immediately felt sick to my stomache and threw up. I went home after my shift and went to bed. After 5 hours, I woke up and went to the couch to lay down while kids watched TV. Next, I woke to several people in my house. I had a seizure and my son had called 911. I was taken to emergency department. I was admitted and stayed 2 nights in the hospital.

VAERS ID: 917122 (history)
Form: Version 2.0

Age: 35.0
Sex: Male
Location: New York
Vaccinated: 2020-12-31
Onset: 2020-12-31
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Blood gases, Blood test, Burning sensation, Chest X-ray, Computerised tomogram, Cough, Dizziness, Headache, Pyrexia, SARS-CoV-2 test, Tremor
SMQs:, Anaphylactic reaction (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Parkinson-like events (broad), Noninfectious encephalopathy/delirium (broad), Vestibular disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Networking, Januvia, amlodopine, losartan
Current Illness: None
Preexisting Conditions: Crohns disease, diabetes, high blood pressure, sleep apnea
Allergies: None
Diagnostic Lab Data: 1/1/2021 covid test, bloodwork, CT scan, chest x ray, bloodgas test
CDC Split Type:

Write-up: A little over ab hour after receiving the vaccine I noticed a burning sensation in my sinuses. By 130am 1/1/2021 I awoke from my sleep terribly dizzy, shaking violently and experiencing a fever of 101.3 F. I took advil and tylenol and fell asleep about an hour later. I woke up with similar symptoms at approximately 830am on 1/1/2021 took an additional dose of advil and tylenol and slept till 12p. I woke up with a bad headache and coughing fits similar to when I had covid back in March. I went to an urgent care who assessed me and ordered me to the ER. medical center administered IV fluids, an inhaler, steroids, epinephrine and benadryl and a few hours later my symptoms had subsided for the most part and a dose of IV antibiotics was administered. I am currently admitted for observation with likely discharge on 1/2/2021.

VAERS ID: 917210 (history)
Form: Version 2.0

Age: 30.0
Sex: Female
Location: Utah
Vaccinated: 2020-12-21
Onset: 2020-12-27
Days after vaccination: 6
Submitted: 0000-00-00
Entered: 2021-01-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / UNK	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood lactate dehydrogenase increased, C-reactive protein increased, Chest X-ray abnormal, Chills, Computerised tomogram thorax, Cough, Diarrhoea, Dyspnoea, Fibrin D dimer normal, Lung consolidation, Lung infiltration, Lung opacity, Procalcitonin increased, Pyrexia, Respiratory viral panel, SARS-CoV-2 test negative, SARS-CoV-2 test positive, Serum ferritin increased, Skin lesion, Sputum culture, Tachycardia, Tachypnoea, White blood cell count increased
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Interstitial lung disease (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Pseudomembranous colitis (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Noninfectious diarrhoea (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (narrow), Infective pneumonia (broad), Dehydration (broad), Sepsis (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: cyclobenzaprine 10mg po tid mirena 52mg IUD
Current Illness: COVID-19 Dec 2 Not hospitalized, mild symptoms
Preexisting Conditions: Acne Depression Anxiety Obesity BMI 35
Allergies: NKDA
Diagnostic Lab Data: see above
CDC Split Type:

Write-up: 30YO F ICU nurse obesity (BMI 35) COVID 19 on Dec 2 symptoms, Dec 3 tested positive for COVID-19. never hospitalized, outpatient only. 12/12 completed isolation 12/21 received vaccine 12/7 developed Fever chills diarrhea SOB cough Urgent care visit. RLL consolidation on CXR given doxycycline 100 mg po bid worse, fever 40 targetoid lesions to LE (started before doxy) WBC 22K tachycardic tachypneic admitted requiring 2-4L oxygen CT

angio without clot, diffuse ground glass and RML dense infiltrate DDimer 7.8 LDH 599 CRP 41 procal 0.67 ferritin 500 Viral respiratory PCR negative Sputum cx with oral flora (pending) COVID ag testing neg COVID PCR 1/3 targets positive (called as indeterminate).

VAERS ID: 917375 (history)
Form: Version 2.0
Age: 60.0
Sex: Female
Location: California
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest X-ray, Chest pain, Dyspnoea, Echocardiogram, Wheezing
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Asthma/bronchospasm (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none
Preexisting Conditions: none
Allergies: environmental allergies
Diagnostic Lab Data: CXR, echocardiogram-January 1, 2021
CDC Split Type:

Write-up: Within 5 minutes of the vaccine, patient had wheezing, shortness of breath and chest pain. patient given epi x 4, decadron, fluids with some improvement and then hospitalized. IN the hospital, patient continued to have chest pain and satting well but with protracted course and is still in the hospital.

VAERS ID: 917497 (history)

Form: Version 2.0
Age: 30.0
Sex: Female
Location: Georgia
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	E20140 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Decreased appetite, Dyspnoea, Hypoaesthesia, Myalgia, Pain in jaw, Paraesthesia, Spinal pain
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Peripheral neuropathy (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Osteonecrosis (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: No known
Diagnostic Lab Data:
CDC Split Type:

Write-up: Vaccine given at 7:05am 12:00noon, 5 hours later, I started experiencing severe chest pain, jaw pain and shortness of breath in which EMS was called and I was taken to the hospital. Since then, I lost feeling in my hands and feet, numbness and tingling. I"ve improved however, during my recovery suffered with spinal pain, shortness of breath, very winded, muscle pain and loss of appetite to especially meat.

VAERS ID: 921188 (history)
Form: Version 2.0

Age: 22.0
Sex: Male
Location: Wisconsin
Vaccinated: 2020-12-17
Onset: 2020-12-21
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2021-01-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Platelet count decreased, Thrombotic thrombocytopenic purpura
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Embolic and thrombotic events, arterial (narrow), Renovascular disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: None known
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Platelet Count, 21-Dec-2020, 2000 cells per microlitre
CDC Split Type:

Write-up: 22 year old patient with no known allergies or medical history admitted 12/21 with TTP and currently being worked up. Currently unclear if related or unrelated to COVID vaccination, but received Pfizer vaccine Thursday 12/17.

VAERS ID: 917702 (history)
Form: Version 2.0
Age: 36.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-24
Onset: 2020-12-29

Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood potassium decreased, Chest discomfort, Cold sweat, Dizziness, Flushing, Hypoaesthesia, Laboratory test normal, Paraesthesia
SMQs:, Anaphylactic reaction (narrow), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Vestibular disorders (broad), Hypersensitivity (broad), Hypoglycaemia (broad), Hypokalaemia (narrow), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vitamin D 2000 iu daily Melatonin 7.5mg daily Mirena (IUD inserted 4/2016)
Current Illness: None
Preexisting Conditions: None
Allergies: Percocet: itching
Diagnostic Lab Data: All tests WNL however my potassium level was low (2.9). Replaced with IV and PO but symptoms still present. Waxing and waning of symptoms.
CDC Split Type:

Write-up: On 12/29: lightheadedness, flushed, felt like I was going to pass out, numbness/tingling down arms and hands, chest tightness, clammy hands and feet.

VAERS ID: 917727 (history)
Form: Version 2.0
Age: 38.0
Sex: Male
Location: New York
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0142 / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Bile duct stone, Biliary dilatation, Blood bilirubin increased, Cholelithiasis, Computerised tomogram abnormal, Full blood count, Headache, Jaundice, Liver function test increased, Metabolic function test, Pain in extremity
SMQs:, Liver related investigations, signs and symptoms (narrow), Cholestasis and jaundice of hepatic origin (narrow), Acute pancreatitis (narrow), Biliary system related investigations, signs and symptoms (narrow), Gallbladder related disorders (narrow), Biliary tract disorders (narrow), Gallstone related disorders (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Folic acid, multivitamin
Current Illness:
Preexisting Conditions: Thalassemia minor, Gilbert"s disease, history of pigment gallstones, hx of splenomegaly
Allergies: None
Diagnostic Lab Data: CT scan, CBC,CMP
CDC Split Type:

Write-up: Patient with a history of thalassemia and Gilbert"s disease, developed severe jaundice three days after vaccination. Had mild headache and sore arm but otherwise felt well. Had labs drawn - found to have highly elevated bilirubin (23) and LFTs in the 700s. Was admitted to the hospital and had CT showing Cholelithiasis, choledocholithiasis and minimal intrahepatic biliary ductal dilatation. Left hospital and was admitted to another facility where plan was for ERCP and cholecystectomy. Ultimately unclear if at all related to the vaccination - may be coincidental.

VAERS ID: 917784 (history)
Form: Version 2.0
Age: 79.0
Sex: Male
Location: Texas
Vaccinated: 2021-01-02
Onset: 2021-01-02
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Brain natriuretic peptide normal, Cardiac arrest, Cardioversion, Coronary arterial stent insertion, Haematocrit decreased, Haemoglobin normal, Intensive care, Red blood cell count decreased, Resuscitation, Syncope, Thirst, Troponin, White blood cell count increased
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Haematopoietic erythropenia (narrow), Haemorrhage laboratory terms (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Embolic and thrombotic events, arterial (narrow), Acute central respiratory depression (broad), Cardiomyopathy (broad), Other ischaemic heart disease (narrow), Hypotonic-hyporesponsive episode (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: WBC 17.3, RBC 4.43, Hct 40.4, Hgb 14.2, Troponin 0.047, BNP 28
CDC Split Type:

Write-up: Pt had vaccination at city site. Waitied 15 min after shot and was cleared to go. Reported to wife that he was very thristy, so they stopped at a convenience store on the way home. While there, he felt worse and asked to go to the Emergency room. They chose Methodist to enter. Pt went to triage and while at triage, had syncopal episode, then full arrest. After short course of CPR and defib, he had ROSC. Was taken to cath lab for intervention (stents) and is now in ICU.

VAERS ID: 917835 (history)
Form: Version 2.0
Age: 18.0
Sex: Male
Location: Minnesota
Vaccinated: 2020-12-30
Onset: 2020-12-31
Days after vaccination: 1
Submitted: 0000-00-00

Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Chest X-ray normal, Chest pain, Culture urine negative, Electrocardiogram PR shortened, Electrocardiogram normal, Headache, Immunoglobulin therapy, Malaise, Pain, Pain in extremity, Paraesthesia, Pyrexia, SARS-CoV-2 antibody test, SARS-CoV-2 test negative, Troponin increased
SMQs:, Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Conduction defects (narrow), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: COVID 19 Positive sept 82020 Few days of malaise headache ,no respiratory symptoms or chest pain Seen By cardiology at College no ECHO or Troponins as no symptoms of chest pain or SOB
Preexisting Conditions: H/o Peri Myocarditis hospitalized 11/2/19-11/5/2019 with ST elevation and elevated Troponins, normal biventricular function treated with IVIG F/u in 2020 doing well also seen in Cardiology Clinic 12/1/2020 Doing well ,no troponins but Holter placed x 24 hrs normal
Allergies: none H/o Cow''s milk hypersensitivity at 1 mos of age ,no problems with milk now
Diagnostic Lab Data: Troponin 4.56 ng/ml after IVIG 2gm/kg 1/2-1/32021 Gammagard 10% (same as 11/2019)
CDC Split Type:

Write-up: Tactile fever ,arm pain, headache and malaise in 24 hrs following injection Next day generalized achiness ,retrosternal chest pain and bilateral forearm tingly pain similar to Nov 2019 and went to Hospital UC,CXR and EKG normal but with short PR interval on EKG ,elevated troponin 3.5 Transferred to hospital troponin 12.1 ng/ml IVIG given SARS IGG positive on admission PCR negative

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VAERS ID: 917855 (history)
Form: Version 2.0
Age: 47.0
Sex: Female
Location: California
Vaccinated: 2020-12-31

Onset: 2021-01-01
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Chest X-ray, Differential white blood cell count, Electrocardiogram, Full blood count, Human chorionic gonadotropin, Metabolic function test, Palpitations, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: NONE. HOWEVER I DID TAKE BENADRYL PROPHYLACTICALLY JUST IN CASE BECAUSE I HEARD OF PEOPLE HAVING ADVERSE REACTIONS TO BOTH MODERNA AND PFIZER COVID VACCINE.
Current Illness: NONE
Preexisting Conditions: NONE
Allergies: SULFA
Diagnostic Lab Data: CBC WITH DIFFERENTIAL, COMPREHENSIVE METABOLIC PANEL, HCG QUANTITATIVE, ECG 12 LEAD, CHEST X-RAY, TYLENOL, IV OF NORMAL SALINE
CDC Split Type:

Write-up: ON 1/1/21 THE DAY AFTER I RECEIVED THE VACCINE I WAS TAKING A NAP AND MY WATCH KEPT SENDING ME ALERTS. HOWEVER I DID NOT CHECK MY WATCH UNTIL 5:30 WHEN I WOKE UP AND I FELT LIKE I WAS HAVING HEART PALPITATIONS. I WENT TO THE EMERGENCY ROOM WHERE I WAS TREATED. I WAS TOLD THAT I WAS FEBRILE WITH A TEMPERATURE OF 102. I WAS GIVEN IV FLUIDS, CHEST X-RAY, EKG AND LAB WORK. I WAS RELEASED ON 1/2/21 AT APPROXIMATELY 0030 (MIDNIGHT).

VAERS ID: 917882 (history)
Form: Version 2.0
Age: 49.0
Sex: Female
Location: New Jersey
Vaccinated: 2020-12-22

Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Dizziness, Dyspnoea, Erythema, Flushing, Nausea, Palpitations, Respiratory distress, SARS-CoV-2 test negative, Swollen tongue, Wheezing
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (narrow), Asthma/bronchospasm (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Biotin 1 mg PO daily, Levothyroxine 112 mcg PO daily, MVI tablet PO daily, Paroxetine 20 mg PO QHS
Current Illness: none reported
Preexisting Conditions: Past Medical History (as noted in H&P) ? Acquired hypothyroidism ? ? Anxiety ? ? Chronic deep vein thrombosis (DVT) of distal vein of lower extremity ? ? Chronic midline low back pain ? ? Complex regional pain syndrome I ? ? Essential hypertension ? ? Fibromyalgia ? ? Gastroesophageal reflux disease without esophagitis ? ? History of pseudoseizure ? ? Laryngospasm ? ? MDD (major depressive disorder) ? ? RSD (reflex sympathetic dystrophy)
Allergies: cefuroxime (shortness of breath), gadolinium (anaphylaxis, March 2018), iodine (anaphylaxis), quinolones (shortness of breath, "paralyze"), gabapentin (hallucinations), vancomycin (itching)
Diagnostic Lab Data: COVID-19 antigen negative, BP 126/52, pulse 96 on admission to ED
CDC Split Type:

Write-up: Patient is hospital employee who completed screening form for COVID-19 vaccine by answering "no" to all contraindication questions. Approx 10 minutes after receiving COVID-19 vaccine dose # 1, patient was still in vaccine clinic area and complained of dizziness, palpitations and flushing. I observed patient fanning herself with papers. She was escorted out of the immediate clinic room, and assessed by paramedics present as having an anaphylactic reaction. Epinephrine 0.3 mg IM and diphenhydramine 50 mg IV given in clinic, Rapid Response was called overhead and patient immediately transported down the hall to the Emergency Dept. In ED, pt was noted as having swollen tongue, large areas of erythema on

face, arms and chest, shortness of breath, nausea, dizziness (per ED physician notes). Pt reported being hospitalized in ICU with COVID disease more than 3 months ago, including intubation (not treated at this hospital), and has been back at work since August 2020. ED physical exam noted bilateral wheezing and patient in acute distress. In ED, pt administered racemic epinephrine 2.25% 0.5 mL via neb, epinephrine 0.3 mg IM, diphenhydramine 50 mg IV, Solu-Medrol 125 mg IV, famotidine 20 mg IV and epinephrine 5 mg/250 mL IV drip (started at 0.118 mcg/kg/min). Acute symptoms reported to resolve in ED. COVID test was negative. ED physician discovered that pt had history of multiple medications, including previous anaphylactic reaction to radiocontrast dye requiring intubation (which was not disclosed on the vaccine screening form). Pt admitted to Telemetry floor for observation. Overnight course was unremarkable, and patient was discharged the following day with prescription for Prednisone taper and prescription for Epi-pen. Advised not to return for second dose of COVID vaccine. EMR updated to reflect possible anaphylactic reaction to Moderna COVID-19 vaccine.

VAERS ID: 917961 (history)
Form: Version 2.0
Age: 54.0
Sex: Male
Location: California
Vaccinated: 2020-12-31
Onset: 2020-12-31
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Angiogram cerebral, Aphasia, Arteriogram carotid, Blood magnesium, Blood thyroid stimulating hormone, Brain natriuretic peptide, C-reactive protein, Chest X-ray, Computerised tomogram head, Culture urine, Differential white blood cell count, Dyspnoea, Dyspnoea exertional, Electrocardiogram, Fibrin D dimer, Full blood count, Glycosylated haemoglobin, Lipids, Magnetic resonance imaging brain, Metabolic function test, Perfusion brain scan, Platelet count, Red blood cell sedimentation rate, Respiratory viral panel, SARS-CoV-2 test, Scan with contrast, Tachycardia, Tachypnoea, Troponin I, Ultrasound kidney, Urine analysis
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Respiratory failure (broad), Infective pneumonia (broad), Dehydration (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes

ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Symbicort inhaler: 2-puffs / twice daily Metoprolol 25mg: 1-tab / daily Lipitor 80mg: 1-tab / daily Zestril 5mg: 1 tab / daily Brilinta 90mg: 1-tab / twice daily Prilosec 20mg: 1-tab / daily HCTZ 25mg: 1-tab / daily ASA 81mg: 1-tab / daily
Current Illness: June 22,2020: COVID19 positive July 2020: Diagnosed with restrictive airway disease, small airway disease, lung nodules November 11, 2020: STEMI, required angioplasty and stent placement
Preexisting Conditions: HTN Conditions highly related to COVID19 infection per provider: Restrictive airway & small airway disease CAD Depression Anxiety
Allergies: Losartan
Diagnostic Lab Data: BASIC METABOLIC Jan 2, 2021 MAGNESIUM Jan 2, 2021 ECG 12-LEAD Jan 1, 2021 D-DIMER,QUANTITATIVE Jan 1, 2021 TROPONIN I Jan 1, 2021 COMPREHENSIVE METABOLIC PANEL Jan 1, 2021 CBC W/ DIFF Jan 1, 2021 C-REACTIVE PROTEIN Jan 1, 2021 US RENAL COMPLETE Jan 1, 2021 MRI BRAIN WO CONTRAST Jan 1, 2021 TROPONIN I Jan 1, 2021 SEDIMENTATION RATE, AUTOMATED Jan 1, 2021 TSH Jan 1, 2021 LIPID PANEL Jan 1, 2021 HEMOGLOBIN A1C Jan 1, 2021 HEMOGRAM 2 (H&H) Jan 1, 2021 PLATELET COUNT Jan 1, 2021 MAGNESIUM Jan 1, 2021 UA W MICROSCOPIC (C&S IF INDICATED) Jan 1, 2021 CT CEREBRAL PERFUSION Dec 31, 2020 CT ANGIOGRAM HEAD NECK W CONTRAST Dec 31, 2020 CT HEAD WO CONTRAST Dec 31, 2020 XR CHEST AP PORTABLE Dec 31, 2020 ECG 12-LEAD Dec 31, 2020 UPPER RESPIRATORY PATHOGEN PANEL W SARS-COV-2 (COVID-19) Dec 31, 2020 CBC W/ DIFF Dec 31, 2020 COMPREHENSIVE METABOLIC PANEL Dec 31, 2020 ER TROPONIN-I Dec 31, 2020 BNP (NT-PROBNP, NT-PRO B-TYPE NATRIURETIC PEPTIDE) Dec 31, 2020 C-REACTIVE PROTEIN Dec 31, 2020 SEDIMENTATION RATE, AUTOMATED Dec 31, 2020
CDC Split Type:

Write-up: I suffer from lingering SOB with exertion after COVID infection. On the night of 12/31/2020 I began to feel more SOB than usual and was unable to correct my SOB with rescue inhaler. Became more SOB with exacerbated tachycardia and tachypnea, My family had to call 911 because I became aphasic and showing signs of possible stroke. I was taken to the ER and admitted for respiratory recovery and to role out stroke. The stroke was ruled out and I recovered with IV prednisone therapy, twice daily and supplemental oxygen. Released after HR, BP, & respiratory effort returned to normal: 01/03/2021

VAERS ID:	918034 (history)
Form:	Version 2.0
Age:	35.0
Sex:	Female
Location:	Michigan
Vaccinated:	2020-12-23
Onset:	2020-12-25
Days after vaccination:	2
Submitted:	0000-00-00
Entered:	2021-01-03

Vaccination / Manufacturer	Lot /	Site / Route

	Dose	
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Exposure during pregnancy, Foetal death, Foetal heart rate abnormal, Foetal hypokinesia, Premature delivery, Stillbirth, Ultrasound Doppler, Ultrasound foetal
SMQs:, Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow), Foetal disorders (narrow), Termination of pregnancy and risk of abortion (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Advair, Pulmocort, levothyroxine, pepcid, prenatal vitamins, singulair, aspirin
Current Illness: None
Preexisting Conditions: Asthma
Allergies: None
Diagnostic Lab Data: Fetal doppler and fetal ultrasound on 12/26/2020
CDC Split Type:

Write-up: I was 28 weeks and 5 days pregnant when I received the first dose of the COVID19 vaccine. Two days later (12/25/2020 in the afternoon), I noticed decreased motion of the baby. The baby was found to not have a heartbeat in the early am on 12/26/2020 and I delivered a 2lb 7oz nonviable female fetus at 29 weeks gestation. I was 35 years old at the time of the fetal demise and the only pregnancy history for this pregnancy included a velamentous cord insertion that was being closely monitored by a high risk OB. My estimated due was March 12, 2021.

VAERS ID: 918051 (history)
Form: Version 2.0
Age: 33.0
Sex: Female
Location: New Jersey
Vaccinated: 2020-12-30
Onset: 2021-01-01
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026LZOA / 2	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Appendicectomy, Appendicitis, Computerised tomogram abdomen abnormal, Laboratory test, Liver function test, Ultrasound abdomen, Ultrasound pelvis
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: prenatal vitamin
Current Illness: n/a
Preexisting Conditions: n/a
Allergies: n/a
Diagnostic Lab Data: Liver panel and chemistry lab: 1/1/21 Abdominal ultrasound: 1/1/21 Abdominal CT scan: 1/1/21 Pelvic Ultrasound: 1/2/21
CDC Split Type:

Write-up: At around 40 hours post vaccination, developed severe abdominal pain and went to an emergency room for evaluation on 1/1/21. Abdominal pain was eventually diagnosed as appendicitis requiring appendectomy on 1/2/21. Emergency room visit and hospital discharged patient early on 1/2/21. It was then determined that the on-call team covering mis-read the CT scan and acute appendicitis was found. Patient then went to Medical Center on 1/2/21 for appendectomy and was discharged later that night following operation.

VAERS ID: 918084 (history)
Form: Version 2.0
Age: 61.0
Sex: Male
Location: California
Vaccinated: 2020-12-21
Onset: 2021-01-03
Days after vaccination: 13
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / IM

Administered by: Military **Purchased by:** ?
Symptoms: COVID-19 pneumonia, Chest X-ray abnormal, Exposure to SARS-CoV-2, Respiratory symptom, SARS-CoV-2 test positive

SMQs:, Cardiomyopathy (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Micardis 40mg daily, Norvasc 10mg daily, Zocor 40 mg daily
Current Illness: None reported
Preexisting Conditions: HTN Hyperlipidemia
Allergies: NKDA
Diagnostic Lab Data: NP swab positive for COVID-19 by PCR on 29 Dec 2020. CXR c/w bilat pneumonia on 03 Jan 2021.
CDC Split Type:

Write-up: Hospitalized with COVID-related pneumonia on 03 Jan 2021. Close contact exposure on 25 Dec, with positive COVID PCR test on 29 Dec... managed as outpatient until respiratory sxms prompted hospitalization on 03 Jan. Care team anticipates at least 4 inpatient days... but patient remains hospitalized at date of this report.

VAERS ID: 918086 (history)
Form: Version 2.0
Age: 31.0
Sex: Male
Location: Unknown
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025120-2A / 1	LA / IM

Administered by: Military **Purchased by:** ?
Symptoms: Electrocardiogram abnormal, Intensive care, Sinus tachycardia, Supraventricular tachycardia
SMQs:, Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations: 2016, Age 27, Flu Vac QS 2016
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: Flu Vaccine
Diagnostic Lab Data: EKG- Intermittent SVT with sinus tachycardia
CDC Split Type:

Write-up: Patient developed SVT 15 minutes after receiving vaccine. Admitted to ICU. ER presentation: BP: 160/109 heart rate 132. No e/o anaphylaxis or allergic reaction.

VAERS ID: 918200 (history)
Form: Version 2.0
Age: 27.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anxiety, Blood chloride increased, Blood phosphorus decreased, Blood potassium decreased, Blood thyroid stimulating hormone normal, Cardiac telemetry normal, Chest X-ray normal, Computerised tomogram thorax normal, Condition aggravated, Dizziness, Echocardiogram normal, Ejection fraction normal, Electrocardiogram ambulatory, Fatigue, Feeling hot, Fibrin D dimer, Flushing, Gait disturbance, Haemoglobin decreased, Pallor, Palpitations, Presyncope, Tachycardia, Troponin normal
SMQs:, Anaphylactic reaction (broad), Haematopoietic erythropenia (broad), Peripheral neuropathy (broad), Haemorrhage laboratory terms (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (broad), Tubulointerstitial diseases (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness: Unknown
Preexisting Conditions: Patient with history of syncope which was attributed to anemia (6-8 episodes in the past). No history of syncope with injections. patient reports increased stress at home and anxiety.
Allergies: None
Diagnostic Lab Data: D- Dimer - 0.5 CT Scan for pulmonary embolism - negative TSH - 2.892 BNP with low potassium (3.3), high chloride (108) and low phos (2.6). All other values within normal limits. Echocardiogram - normal with normal EF Chest xray - normal Troponin negative x3 Hgb 11.7
CDC Split Type:

Write-up: Pfizer-BioNTech COVID-19 Vaccine EUA - Patient witnessed another patient with syncope prior to her injection. She was already anxious about receiving vaccination and this increased her anxiety, though she proceeded with immunization. Patient was in 15 min observation window in a chair and began to feel light-headed like she may pass out. A SWAT was called. With RN assistance, patient was lowered to the floor, with no loss of consciousness. Patient was pale and reported anxiety, racing /pounding heart, and felt hot with facial flushing. Patient was transferred to ED and was noted to be tachycardic (120s), but dropped to 80s. She noted that this episode felt different than her prior syncopal episodes associated with anemia. Patient was observed for 5 hours and discharged to home. Patient returned to ED roughly 2.5 hours later complaining of continued dizziness and unsteady gate. Patient was pale and anxious. Patient reported had not eaten/drank enough during her shift and received vaccine immediately post a stressful shift. Additionally, patient witnessed another patient have syncopal episode prior to her receiving her vaccine which made her anxious. Patient was given IV fluids and had electrolytes replacement. Patient additionally received diazepam. Patient was discharged at 2358 on 12/23. Patient returned to ED on 12/24 at 0239 complaining of near syncope and lightheadedness. Patient had tachycardia and self-reported palpitations. Received IV fluids and observation on telemetry with no rhythm disturbance. Patient discharged 1428 on 12/24. On 12/29, patient returned to ED at 0326 for continued dizziness, fatigue and near syncope. Was admitted for cardiac evaluation. Noted to have unprovoked tachycardia and was discharged with a Halter Monitor to evaluate cardiac symptoms. patient was discharged 12/31 at 1619

VAERS ID:	918211 (history)
Form:	Version 2.0
Age:	52.0
Sex:	Female
Location:	Florida
Vaccinated:	2020-12-31
Onset:	2020-12-31
Days after vaccination:	0
Submitted:	0000-00-00
Entered:	2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dysphonia, Eyelids pruritus, Intensive care, Paraesthesia oral, Pruritus, Rash, Rash maculo-papular, Swelling face
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Periorbital and eyelid disorders (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness: Unknown
Preexisting Conditions: Asthma, OSA, history of anaphylaxis, GERD
Allergies: Benzonatate - brochospasm and laryngospasm requiring intubation NSAIDS - petichiae Keflex - rash - tolerates cefepime Lanolin - rash Septra - rash
Diagnostic Lab Data: benadryl multiple rounds of epinephrine methylprednisoolone famotidine
CDC Split Type:

Write-up: Pfizer-BioNTech COVID-19 Vaccine EUA - Patient with history of anaphylaxis requiring intubation to benzonatate. Patient answered "no" to questionnaire about allergic reactions prior to vaccination. 11 minutes after vaccination, patient reported tingling of lips and swelling of face. Developed hoarseness. SWAT was called and patient given benadryl and taken to ED (1055). Patient received steroids and H1/H2 blockers in addition to epinephrine. Patient brought to ICU for monitoring. Patient continued on therapy and was discharged 1/2 at 1113. Patient returned to ED on 1/3 at 1558 with macular papular rash on leg, chest and back with itching on eyelids and face. No respiratory involvement. Patient given benadryl and predinisone and discharged from ED at 2016.

VAERS ID: 918409 (history)
Form: Version 2.0
Age: 38.0
Sex: Male
Location: California
Vaccinated: 2020-12-18
Onset: 2020-12-25
Days after vaccination: 7

Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Cough, Dyspnoea, Fatigue, Intensive care, SARS-CoV-2 test positive, Sinus tachycardia
SMQs:, Anaphylactic reaction (broad), Supraventricular tachyarrhythmias (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Infective pneumonia (broad), Dehydration (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 7 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: no known allergies
Diagnostic Lab Data: 12/24: SARs CoV 2 RNA, RT PCR (+)
CDC Split Type:

Write-up: Patient complained of increased shortness of breath, generalized weakness and fatigue with mild cough worsening today and was admitted on 12/25/20. Patient is an employee of the hospital in the ICU and received the covid-19 vaccine on 12/25/20. Patient believes symptoms started after the vaccination. On admission, patient was in sinus tachycardia with O2 saturation 91% on room air. Tested SARs CoV 2 RNA, RT PCR positive on 12/24/20. Transferred to ICU for closer monitoring after transitioning to high flow nasal cannula on 12/26. Patient recovered and discharged on 12/31/20.

VAERS ID: 918552 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-30
Onset: 2021-01-01
Days after vaccination: 2
Submitted: 0000-00-00

Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 5	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chills, Diarrhoea, Influenza virus test negative, Nausea, Pyrexia, SARS-CoV-2 test negative
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: Lupus
Preexisting Conditions: Lupus
Allergies: NKA
Diagnostic Lab Data: Pt was tested for Influenza and COVID and lab work was normal
CDC Split Type:

Write-up: Severe diarrhea, cold chills, 101 fever, and nausea

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VAERS ID: 918608 (history)
Form: Version 2.0
Age: 78.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, Hyperhidrosis, Injection site cellulitis, Injection site erythema, Injection site pain, Injection site swelling, Injection site warmth, Pyrexia, Retching, SARS-CoV-2 test negative

SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: OTC: vitamin D, cranberry caps, polycarbophil, senokot, duloxetine, estradiol VC, furosemide, metoprolol, mirabegron ER, rivaroxaban
Current Illness: none
Preexisting Conditions: HTN, T2DM, dementia, Hx PE/DVT
Allergies: codeine-N/V, darvan/sulfa drugs- no reaction documented
Diagnostic Lab Data: Rapid COVID-19 test on admission- Negative was COVID positive in July 2020
CDC Split Type:

Write-up: "Pfizer-BioNTech COVID-19 Vaccine" 12/29 patient developed SOB, fever tmax 103 degrees F, diaphoretic, dry heaves all started approximately 16 hours after vaccination given. patient then transferred to Hospital for further treatment and observation. 12/30 seen at injection site- erythema, swelling, warmth and tenderness Discharged back to home on 1/1 with RX for cephalexin to treat cellulitis of injection site

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VAERS ID: 918740 (history)
Form: Version 2.0
Age: 36.0
Sex: Unknown
Location: Unknown
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Lip pruritus, Throat irritation

SMQs:, Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Food allergy (known allergies: Shrimp)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020508654

Write-up: Scratchy throat; itching lips; This is a spontaneous report from a non-contactable nurse. A 36-year-old patient of an unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), lot number EK5730, via an unspecified route of administration on Right arm from 22Dec2020 to 22Dec2020 as single dose for COVID-19 immunization. Medical history included food allergy (Shrimp). The patient''s concomitant medications were not reported. The patient experienced scratchy throat and itching lips for approximately 1.5 hours starting 20 mins post vaccine on 22Dec2020. The event caused prolonged hospitalization. The outcome of the events was recovered on 22Dec2020. No follow-up attempts are possible. No further information is expected.; Sender''s Comments: Based on temporal association, a possible contributory role of suspect BNT162B2 vaccine cannot be excluded for reported events throat irritation and lip itching. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 918741 (history)
Form: Version 2.0
Age: 74.0
Sex: Male
Location: South Carolina
Vaccinated: 2020-12-15
Onset: 2020-12-15
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Ataxia, Chills, Confusional state, Decreased appetite, Delirium, Fatigue, Headache, Hypoxia, Incontinence, Myalgia, Pyrexia, SARS-CoV-2 test
SMQs:, Rhabdomyolysis/myopathy (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (narrow), Dementia (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (narrow), Noninfectious meningitis (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Dehydration (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Diabetes mellitus
Allergies:
Diagnostic Lab Data: Test Name: tested for COVID-19 via nasal swab; Result Unstructured Data: Test Result:Unknown Result
CDC Split Type: USPFIZER INC2020509173

Write-up: delirious; hypoxic; fever; ataxic; incontinent; confused; Chills; HA; anorexia/had no appetite; myalgias; extreme fatigue; slept all and had no appetite; This is a spontaneous report from a contactable physician. A 74-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 15Dec2020 16:00 at single dose for COVID-19 immunization. Medical history included diabetes mellitus (DM). The patient has no known allergies. The patient''s concomitant medications were not reported. He was an ER doctor and the medical director of his hospital. The patient was asymptomatic when he got the vaccine on 15Dec2020. 3 hours after the vaccine he began to get chills, HA, anorexia, myalgias, and extreme fatigue. This worsened and he slept all and had no appetite. On 19Dec2020 he woke up delirious with a fever and was ataxic, hypoxic, incontinent, and confused. The patient was hospitalized due to the events on 15Dec2020. The events also caused prolonged hospitalization due to the events. The patient was not diagnosed with COVID prior to vaccination. The patient was tested for COVID via nasal swab post vaccination with unknown results. The patient did not receive any other vaccines within 4 weeks prior to COVID vaccine. The outcome of the events was not recovered. Therapeutic measures were taken as a result of the events as the patient required oxygen, plasma, and remdisivir. The batch/lot numbers for the vaccine, BNT162B2, were not provided and will be requested during follow up.; Sender''s Comments: A possible causal association between administration of BNT162B2 and the onset of reported serious events might not be excluded, considering the plausible temporal relationship. Fever, chills, headache, fatigue and muscle pain are the known adverse event profile of the suspect product. More information such as

detailed underlying medical conditions and concomitant medications are needed for fully medical assessment. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 918758 (history)
Form: Version 2.0
Age: 61.0
Sex: Female
Location: New Hampshire
Vaccinated: 2020-12-21
Onset: 2020-12-22
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / OT

Administered by: Private **Purchased by:** ?
Symptoms: Lymphadenopathy, SARS-CoV-2 test negative, Small intestinal obstruction
SMQs:, Gastrointestinal obstruction (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; COLESTID; STELARA; B12 [CYANOCOBALAMIN]
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Crohn"s disease
Allergies:
Diagnostic Lab Data: Test Date: 20201223; Test Name: Nasal Swab; Test Result: Negative
CDC Split Type: USPFIZER INC2020512697

Write-up: Small bowel obstruction with diffuse bowel and pelvic lymphadenopathy 36 hours after injection; Small bowel obstruction with diffuse bowel and pelvic lymphadenopathy 36 hours after injection; This is a spontaneous report from a contactable physician (patient). A 61-years-old female patient (not pregnant) received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number: EH9899), intramuscularly on 21Dec2020 12:30 to at single dose on left arm for COVID-19 immunization in hospital. Medical history

included crohn"s disease. No known allergies. Concomitant medications within 2 weeks of vaccination included estradiol, progesterone, colestipol hydrochloride (COLESTID), ustekinumab (STELARA), cyanocobalamin (B12). The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient experienced small bowel obstruction with diffuse bowel and pelvic lymphadenopathy 36 hours after injection on 22Dec2020 22:00 with outcome of recovered in Dec2020. The adverse events resulted in emergency room/department or urgent care, hospitalization for 3 days. Therapeutic measures were taken as a result of event included inpatient observation, nothing by mouth (reported as NPO), intravenous fluids. No COVID prior vaccination, COVID test nasal swab was negative on 23Dec2020 post vaccination. It was not reported as serious.; Sender"s Comments: There is not a reasonable possibility that reported events small bowel obstruction and lymphadenopathy are related to BNT162B2 vaccine. The patient had underlying Crohn"s disease, which put patient at risk of developing the event. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 918763 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Unevaluable event
SMQs:

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Guillain Barre syndrome

(Caller had Guillain Barre in 2011 at age 65)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020513681

Write-up: She was hospitalized for a month Occurred after a flu two months prior and a stomach flu 2-3 weeks prior; She was hospitalized for a month Occurred after a flu two months prior and a stomach flu 2-3 weeks prior; This is a spontaneous report from a contactable consumer. A 75-year-old female patient received bnt162b2 (BNT162B2, lot number and expiry date were not reported), via an unspecified route of administration on an unspecified date at a single dose for covid-19 vaccination. Medical history included guillain-barre syndrome from 2011 at the age of 65 and Levaquin allergy. The patient''s concomitant medications were not reported. On an unspecified date, the patient was hospitalized for a month that occurred after a flu two months prior and a stomach flu 2-3 weeks prior. The outcome of the events was unknown. Information about lot/batch number has been requested.

VAERS ID: 918770 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: Georgia
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	ELO140 / 1	LA / OT

Administered by: Work **Purchased by:** ?
Symptoms: Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020514065

Write-up: Difficulty breathing; This is a spontaneous report from a contactable pharmacist. A 54-year-old female patient received her first dose of intramuscular BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 22Dec2020 at 04:15 PM at single dose in left arm for COVID-19 immunisation at the age of 54-year-old. Lot number was ELO140. Medical history was unknown, concomitant medications were unspecified. Patient was not pregnant at the time of vaccination. On 22Dec2020 at 04:30 PM, the patient experienced difficulty in breathing, and she was hospitalized for one day. The patient was treated with EPI for the event. The patient was recovering from the event.; Sender"s Comments: Based on the compatible temporal association, the Company considers the event difficulty in breathing is possibly related to vaccination with BNT162B2. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 918774 (history)
Form: Version 2.0
Age: 93.0
Sex: Male
Location: Oregon
Vaccinated: 2020-12-22
Onset: 2020-12-01
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Abdominal pain, Blood pressure increased, Hypertension, Nausea
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypertension (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions: Medical History/Concurrent Conditions: Diabetic; Irritable bowel
Allergies:
Diagnostic Lab Data: Test Date: 202012; Test Name: blood pressure; Result Unstructured Data: Test Result:high
CDC Split Type: USPFIZER INC2020514106

Write-up: abdominal pain; nausea; high blood pressure; This is a spontaneous report from a contactable consumer. A 93-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL0140), via an unspecified route of administration on 22Dec2020 at a single dose for COVID-19 immunization. Medical history included diabetic and irritable bowel. The patient''s concomitant medications were not reported. The patient received the COVID vaccine and had abdominal pain, nausea and high blood pressure within 12 to 18 hours of vaccine received. The events lead to nursing home to emergency room and admitted to hospital. The patient was hospitalized due to events since 23Dec2020. Outcome of the events was recovering.

VAERS ID: 918783 (history)
Form: Version 2.0
Age: 29.0
Sex: Male
Location: Texas
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Aphasia, Balance disorder, Cardiac assistance device user, Cardiac disorder, Cerebrovascular accident, Computerised tomogram, Dyskinesia, Dysphemia, Echocardiogram, Ejection fraction decreased, Hemiparesis, Hypoaesthesia, Impaired work ability, Magnetic resonance imaging, Muscle fatigue, Neurological examination abnormal, Occupational exposure to SARS-CoV-2, Paraesthesia, Speech disorder, Ventricular tachycardia
SMQs:, Torsade de pointes/QT prolongation (narrow), Rhabdomyolysis/myopathy (broad), Cardiac failure (narrow), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Ventricular tachyarrhythmias (narrow), Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Torsade de pointes, shock-associated conditions (narrow), Dementia (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Dyskinesia (narrow), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow), Vestibular disorders (broad), COVID-19 (narrow), Sexual dysfunction

(broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Name: CT; Result Unstructured Data: Test Result:Unknown result; Test Name: Echocardiogram; Result Unstructured Data: Test Result:Unknown result; Test Name: Ejection fraction; Test Result: 25 %; Test Name: MRIs; Result Unstructured Data: Test Result:Unknown result
CDC Split Type: USPFIZER INC2020516120

Write-up: left sided weakness; it has weakened his heart; stutter; severe stroke like symptoms; Ventricular tachycardia/help keep his heart rate at bay; Loss of balance; extreme numbness and tingling in left hand and foot; tingling in left hand and foot; oral motor impairment; mouth weakness and not coordinated/mouth is fatigued easily; Issues finding words and trouble speaking; Issues finding words and trouble speaking; Ejection fraction down to 25%; The initial case was missing the following minimum criteria: adverse event. Upon receipt of follow-up information on 28Dec2020, this case now contains all required information to be considered valid. This is a spontaneous report from a contactable other healthcare professional (HCP). A 29-years-old male patient received bnt162b2 (lot number: EJ1685), intramuscular (deltoid left) on 21Dec2020 at 05:30 at 0.3 mL single (first dose) for Covid. Medical history was reported as "none". Concomitant medications were not reported. The patient previously received Flu vaccine in Oct2020 for immunization. The patient is an Occupational Therapist, and he called to report an adverse event that he experienced with the first dose of the COVID Vaccine. He received the vaccine last Monday, 21Dec2020 at 5:30AM before his shift at work, then 20 minutes later, he was having severe stroke like symptoms. He experienced severe left sided weakness, loss of balance, extreme numbness and tingling in his left hand and foot, he had issues finding his words and he couldn"t speak, and he had an oral motor impairment where his mouth was weak and not coordinated. The staff at the hospital did a neurological exam on him, and he failed, so he had to go to the emergency room (ER). The patient added that he was already in the hospital when this happened, and the ER doctors suspected that he had a CVA, and they gave him TPA to prevent any permanent brain damage and it worked. The patient then added that due to the shock of this whole event, from everything that happened, it has weakened his heart. Reportedly, he is a healthy 29 year old man, with no preexisting conditions, and he works out, and he has no heart conditions, but he had to get a cardiology follow up a few days after he got the vaccine, because he started going in to Ventricular Tachycardia, which he had never had in his life. So, the doctors at the hospital went ahead and did an Echocardiogram and an EKG, and he was told that his Ejection Fraction is down to 25%. He stated his heart is so weak, that he cannot work right now, but the structure of his heart is fine and has not had any damage. The hospital staff thought that maybe the patient had a chronic heart issue that he just did not know about,

and that the stress of this event maybe made it kick into overdrive, but he states that the cardiologist said that was not the case, because the structure of his heart is fine, and the only thing they can see is that the heart is pumping weak. One physician even suggested that due to the shock of the event, he might have Takotsubo Cardiomyopathy, which is a broken heart, but because the structure of his heart is okay, it should be reversible. He stated that he is hoping he will heal up good, because he is young and has no pre-existing conditions. He added that his heart is in such a state right now; he has to wear an external defibrillator. The patient stated that all these happened about 20 minutes after he received the vaccine, and he was admitted to the hospital from 21Dec2020 to 25Dec2020. His neurological symptoms have resolved except that he has a stutter that he did not have before and his mouth is fatigued easily, so he has to slow down when he is eating, but now he can eat regular for the most part. The patient confirmed that he was not specifically prescribed the product; it was administered to him at his place of work, but it was optional. He stated he considered how he is working with COVID patients every day, and given the circumstance, he thought that it would be a best practice for him to get the vaccine. He had not gone to his primary care doctor in a while because he had been fine and healthy, but he called them and found out that his primary care had retired, so he has to find a new one now. Regarding the issues finding words and trouble speaking, he stated that he has improved, but it is still ongoing, he is just stuck in a plateau zone. With the Ventricular Tachycardia, he stated that this is an ongoing issue, as he has to wear the life vest even though he has no need to activate it yet. He did have one minor bought of the VTach, but because he is a therapist, he knows how to take care of it with relaxation techniques, he knows how to manage it. He had one bought of VTach the evening prior, but he was able to get it under control. The doctors have him on medication to help keep his heart rate at bay. He has never had to use medication before and is on the following medications to help keep his heart rate at bay: Metoprolol 25mg one tablet once daily by mouth and Lisinopril 5mg one tablet once daily by mouth. The VTach has improved, it was good enough he was able to discharge home, but it is still a concern. His cardiologist said that, basically his hope, is that once his body recover from the whole shock of everything, then his ejection fraction will heal, and his heart will heal. He again stated that the doctor told him that the structure of his heart is perfectly fine; he has thick walls in his heart, no leaking valves, and the heart was not conducting any abnormal signals. The doctor just said that right now, his heart is super weak and that it is an acute problem. With the Takotsubo Cardiomyopathy, he states that two doctors mentioned this diagnosis, but he confirmed that he was not actually diagnosed with this issue, he was just diagnosed with Ventricular Tachycardia. The outcome of the ejection fraction down to 25% was unknown to the patient at this time as he has not had another EKG or echocardiogram, but the cardiologist told him that the cardiologist expects that this will not be resolved quickly anyway. The patient confirmed that he did not receive any other vaccines on the same day he received the COVID vaccine. The only other vaccine he had this year was the flu vaccine which he got back in Oct2020. He has gone on to his online portal and there are the bloodwork results and all the imaging results on there from his CTs and MRIs, but he did not see the EKG or Echocardiogram results yet. He does not have this pulled up at this time, but he does have access to this stuff and can provide it later, if requested. He is curious about the next steps from here to how his case is processed. He is also curious if this information would help Pfizer make modifications to the vaccine if it is found that a lot of people are having the same reaction as he did. He is also wondering, given his situation, that probably he is not going to get the second dose, for his safety, but he is wondering what percentage of effectiveness the first dose does having just covered. The events left sided weakness, loss of balance, extreme numbness and tingling in left hand and foot resolved on 25Dec2020; severe stroke like symptoms and oral motor impairment; mouth weakness and not coordinated/mouth is fatigued easily resolved in 2020. The events ventricular tachycardia/help keep his heart rate at bay and issues finding words and trouble speaking were resolving, stutter had not resolved while the outcome of the events it has weakened his heart, and ejection fraction down to 25% was unknown.; Sender"s Comments: The reported information is unclear and does not allowa meaningful assessment

of the case. It will be reassessed upon receipt of follow up information. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 918789 (history)
Form: Version 2.0
Age: 30.0
Sex: Female
Location: Massachusetts
Vaccinated: 2020-12-18
Onset: 2020-12-01
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Anxiety, Blood pressure increased, Echocardiogram, Heart rate increased, Immediate post-injection reaction, Myocardial necrosis marker increased, Palpitations
SMQs:, Neuroleptic malignant syndrome (broad), Myocardial infarction (broad), Arrhythmia related investigations, signs and symptoms (broad), Hypertension (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201218; Test Name: blood pressure; Result Unstructured Data: Test Result:elevated; Test Date: 202012; Test Name: echocardiogram; Result Unstructured Data: Test Result:Unknown results; Test Date: 202012; Test Name: heart rate; Result Unstructured Data: Test Result:racing; Test Date: 20201218; Test Name: heart rate; Result Unstructured Data: Test Result:accelerated; Test Date: 202012; Test Name: heart enzymes; Result Unstructured Data: Test Result:elevated
CDC Split Type: USPFIZER INC2020516465

Write-up: She had an immediate reaction of accelerated heart rate and elevated blood pressure; She has very slightly elevated heart enzymes.; She had an immediate reaction of accelerated heart rate and elevated blood pressure; She"s very anxious and very anxious tonight being alone at the hospital.; This is a spontaneous report from a contactable consumer (patient"s father). A 30-years-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 18Dec2020 at single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient had an immediate reaction of accelerated heart rate and elevated blood pressure. She was monitored for an hour and it subsided. On unspecified date in Dec2020 at 1:00am in the morning she had racing heart rate and went to ER. She was still in hospital. Things were not entirely stabilized. She had very slightly elevated heart enzymes in Dec2020 with outcome of unknown. They keep her overnight for an echocardiogram in morning. The patient was very anxious being alone at the hospital. Caller questioned if this has been reported with the vaccine. Lot/Batch and Expiry date has been requested.

VAERS ID: 919036 (history)
Form: Version 2.0
Age: 89.0
Sex: Male
Location: Indiana
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Bladder catheter replacement, Dehydration, Hyperhidrosis, Loss of consciousness, Nausea, Speech disorder, Vomiting
SMQs:, Torsade de pointes/QT prolongation (broad), Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypoglycaemia (broad), Dehydration (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness: unknown
Preexisting Conditions: Unknown
Allergies: Sulfa antibiotics
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient with extreme nausea and vomiting that started soon after receiving the Moderna vaccine. Patient with loss of consciousness, diaphoresis and garbled speech during a foley catheter exchange thought to be from dehydration. Patient was admitted to Hospital for observation for 2 days

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VAERS ID: 919087 (history)
Form: Version 2.0
Age: 50.0
Sex: Male
Location: California
Vaccinated: 2020-12-23
Onset: 2020-12-27
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Antinuclear antibody negative, C-reactive protein increased, Fibrin D dimer increased, Hepatitis C antibody negative, Pericarditis, Red blood cell sedimentation rate normal, Troponin T increased
SMQs:, Haemorrhage laboratory terms (broad), Systemic lupus erythematosus (broad), Myocardial infarction (narrow), Chronic kidney disease (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Aspirin daily
Current Illness: No illness one month prior to or at time of vaccination

Preexisting Conditions: None
Allergies: NKDA
Diagnostic Lab Data: 12/29 ANA negative. CRP 7.2 mg/dL. ESR 8 mm/hr. 12/28 Troponin T 67 ng/L. Hep C Ab = Non reactive. D-dimer 319 ng/mL.
CDC Split Type:

Write-up: Acute Pericarditis. Patient was admitted from 12/27-12/28/2020 at hospital by cardiology team who strongly felt the acute pericarditis was due to the Pfizer Vaccine (Dr. was senior cardiologist).

VAERS ID: 919129 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Hawaii
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood test, Computerised tomogram, Hypoaesthesia, Magnetic resonance imaging, Paraesthesia
SMQs:, Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Losortan HCTZ, Claritin
Current Illness: no
Preexisting Conditions: Hypertension
Allergies: Vicodin
Diagnostic Lab Data: Blood work, CT, MRI to rule out the possibility of a stroke
CDC Split Type: vsafe

Write-up: After the 15 min monitoring, I went back to work 15 min later. the left side of my face started tingling which went to a numbing feeling down the left side of my body affecting

my neck, shoulder, arm, elbow and up the upper left torso. My face and neck was numb for about 48 hrs and the rest came back to sensation within 24 hrs. D/T the numbness I was admitted into the hospital for a 24hr observation.

VAERS ID: 919152 (history)
Form: Version 2.0
Age: 41.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-21
Onset: 2020-12-27
Days after vaccination: 6
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Appendicectomy
SMQs:

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: "Pfizer-BioNTech COVID-19 Vaccine EUA." Patient received first dose of vaccine on 12/21/2020. Patient called on 1/3/2021 to notify that she developed symptoms on 12/27/2020 and had an appendectomy on 12/28/2020 at another facility. Patient also reports that CDC V-Safe Application was used to report event as well. Patient reports that she is recovering well.

VAERS ID: 919154 (history)

Form: Version 2.0
Age: 38.0
Sex: Female
Location: Iowa
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram normal, Ear pain, Facial paralysis, Hypoaesthesia, Hypoaesthesia oral, Pain in extremity, Paraesthesia
SMQs:, Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Hearing impairment (broad), Tendinopathies and ligament disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Probiotic
Current Illness: N/A
Preexisting Conditions: Crohns Disease
Allergies: Azithromycin (patient has an upset stomach when taking), patient also has sensitivity when eating chocolate, pineapple or highly processed foods (due to Crohn"s Disease)
Diagnostic Lab Data: CT scan done on 1-2-21 -negative for hemorrhage
CDC Split Type:

Write-up: Patients adverse reactions started day of vaccination with right arm pain up to right ear as well as complete tongue numbness. On 1-1-21 patient had increased Bell"s Palsy symptoms including; inability to raise left eyebrow, inability to close left eye in its entirety, teeth being numb on left side, and numbness and tingling in left foot and left hand (from palm to fingers). ER physician prescribed on 1-1 Prednisone, Keflex and Valtrex. Patient went to ER again on 1-2-21 with lower extremity numbness on left side that is moving proximally toward her hip. Patient went home on 1-3-21 with an RX for Prednisone as well as Valtrex. Symptoms have improved but have not fully resolved at this time.

VAERS ID: 919230 (history)

Form: Version 2.0
Age: 41.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-18
Onset: 2020-12-26
Days after vaccination: 8
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ 1685 / UNK	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: SARS-CoV-2 test positive
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: no
Diagnostic Lab Data:
CDC Split Type:

Write-up: She was hospitalized on 01/04 but exact situation unknown. COVID +. Hospitalized at Medical Center.

VAERS ID: 919252 (history)
Form: Version 2.0
Age: 27.0
Sex: Female
Location: Maine
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest discomfort, Headache, Palpitations, Pyrexia, Rash, Skin discomfort
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: NKA
Diagnostic Lab Data: EE was seen in the emergency room and admitted to Hospital. She remains inpatient today 1/4/21.
CDC Split Type:

Write-up: Employee received COVID 19 vaccination at 9:45am on 12/30/20. ~15 min. later she developed a rash down her left arm, then down her Rt. arm. about 4 hours later she decided to go to the emergency room for Hearty Palpitations, Fever, Chest discomfort and feeling of generalized sunburn. Later developed severe headache..

VAERS ID: 919320 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	RA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Blood test, Dizziness, Hyperhidrosis, Immediate post-injection reaction, Peripheral swelling, SARS-CoV-2 test
SMQs:, Cardiac failure (broad), Angioedema (broad), Neuroleptic malignant syndrome

(broad), Anticholinergic syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Vestibular disorders (broad), Hypersensitivity (narrow), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Toperol (high blood pressure) Adderall - 20mg Calcitrol- .5mcg Soduim bycar-650mg
Current Illness: Stage 5 Kidney failure
Preexisting Conditions: Stage 5 Kidney failure
Allergies: Mushrooms Bactriun
Diagnostic Lab Data: Blood test, covid test
CDC Split Type: vsafe

Write-up: Right arm swelling very bad right after shot, next day woke up to get ready to work I started to get light headed, dizzy, sweating, felt like I was going to pass out. My husband then called 911. They took me to Hospital ,I stayed there for a couple hours then released. They told me to stay home and the next day I felt fine. I did a televisit with my Nephrologist (Kidney doctor) the following week.

VAERS ID: 919428 (history)
Form: Version 2.0
Age: 58.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-26
Onset: 2020-12-30
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram, Burning sensation, Catheterisation cardiac normal, Chest pain, Discomfort, Feeling hot, Myocardial infarction, Pain in extremity, Troponin increased
SMQs:, Peripheral neuropathy (broad), Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No

Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: clopidogrel lisinoprel carvedilol rosuvastatin aspirin symbicort melatonin mvi vitamin d3
Current Illness: none
Preexisting Conditions: mitral valve takotsubo cardiomyopathy myocardial infarction 2011 migraines htn
Allergies: nka
Diagnostic Lab Data: Angiogram 12/31/2020 Troponin levels 12/30/2020 <0.02; 12/31/2020 1.7 PER PATIENT
CDC Split Type:

Write-up: Felt slight warmth throughout body about 5 minutes after vaccine. Disappeared 2 minutes later. Arm started to feel sore as the day went on and was very sore by nighttime. Next day, arm started to feel better and over the next 3 days was no longer sore. On the morning of the 30th, woke up feeling fine, took a 3.5 mile walk and felt fine. Around 12:30 pm, experienced sudden pain and a burning sensation in the chest and both upper arms. Thought it was possibly heartburn ; took a Prilosec. The discomfort (mild but steady) continued so checked blood pressure which was 141/91. Called cardiologist and went to emergency room per instruction, around 1:30. Admitted overnight with the diagnosis of a mild heart attack and performed a heart catheterization where they found no major blockage. One artery noted 30% blocked but that overall heart function looked good. Discharged on 12/31/2020. -reported by patient via email, on 1/3/2021 @ 4:49pm

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VAERS ID: 919436 (history)
Form: Version 2.0
Age: 29.0
Sex: Male
Location: Illinois
Vaccinated: 2020-12-29
Onset: 2021-01-01
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Blood test, Computerised tomogram, Dysphagia, Eyelid function disorder, Facial paralysis, Hypoaesthesia, Magnetic resonance imaging
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome

(broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hearing impairment (broad), Periorbital and eyelid disorders (narrow), Ocular motility disorders (narrow), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Strained neck resulting in numbness in extremities. This and facial numbness prompted me to see my PCP who urged me to go to the ER immediately. I went to Medical Center''s ED, after which I was admitted to the neuro ICU for a possible brain bleed, which after an MRI was determined to be a cavernoma.
Preexisting Conditions: Depression, anxiety, seasonal allergies
Allergies:
Diagnostic Lab Data: Full bloodwork, CT, MRI as part of inpatient hospitalization 1/2/2021-1/3/2021 at Medical Center.
CDC Split Type:

Write-up: Bell?s palsy, right side of face is numb, with difficulty closing eyes, smiling, raising eyebrows, eating, drinking, swallowing.

VAERS ID: 919546 (history)
Form: Version 2.0
Age: 51.0
Sex: Female
Location: Alabama
Vaccinated: 2020-12-29
Onset: 2021-01-01
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Antinuclear antibody positive, Antiphospholipid antibodies, Aphasia, Blood cholesterol increased, Blood homocysteine, Coagulation time, Craniotomy, Low density lipoprotein increased, Mechanical ventilation, Thrombotic stroke
SMQs:, Dyslipidaemia (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Systemic lupus erythematosus (narrow), Ischaemic central nervous system vascular conditions (narrow), Dementia (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Acute central respiratory depression

(broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Lipodystrophy (broad), Respiratory failure (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Plaquenil Fioricet Midrin
Current Illness: PMH- Lupus; HTN; marked hyperlipidemia
Preexisting Conditions: Lupus Hypertension Hyperlipidemia
Allergies: None listed
Diagnostic Lab Data: Total cholesterol- 465 LDL- 351 ANA (+) Labs pending- Lupus AC; antiphospholipid AB, homocysteine levels; hypercoagulability profile
CDC Split Type:

Write-up: thrombotic stroke -necessitating hospitalization; and craniotomy; required mechanical ventilator for 2 days. Patient now extubated, breathing on her own. Patient remains hospitalized with marked deficits (aphasic)

VAERS ID: 919584 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: Unknown
Vaccinated: 2021-01-03
Onset: 2021-01-03
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Chest discomfort, Dizziness, Throat tightness
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Vestibular disorders (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness:
Preexisting Conditions: none
Allergies: azithromycin
Diagnostic Lab Data:
CDC Split Type:

Write-up: Lightheadedness, throat tightness. Increasing chest tightness. History of atrial fibrillation and bilateral breast implants. Received two doses of epinephrine and one dose of diphenhydramine.

VAERS ID: 919593 (history)
Form: Version 2.0
Age: 64.0
Sex: Male
Location: Iowa
Vaccinated: 2020-12-30
Onset: 2021-01-02
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	RA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Arthritis bacterial, Culture wound positive
SMQs:, Arthritis (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: none
Preexisting Conditions: hypertension, sleep apnea, osteoarthritis
Allergies: opioids
Diagnostic Lab Data: Positive wound cultures.

CDC Split Type:

Write-up: Patient developed a septic knee (history of arthroplasty) need for immediate surgery, hospitalization and months to years of antibiotics in his future now.

VAERS ID: 919604 (history)
Form: Version 2.0
Age: 30.0
Sex: Female
Location: Mississippi
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Alanine aminotransferase normal, Anion gap decreased, Aspartate aminotransferase normal, Basophil count normal, Basophil percentage decreased, Blood albumin normal, Blood alkaline phosphatase normal, Blood bilirubin normal, Blood calcium normal, Blood chloride normal, Blood creatinine normal, Blood glucose normal, Blood magnesium normal, Blood potassium decreased, Blood sodium normal, Blood thyroid stimulating hormone normal, Blood urea nitrogen/creatinine ratio, Blood urea normal, Carbon dioxide decreased, Chest X-ray normal, Chest discomfort, Cough, Dizziness, Eosinophil count decreased, Eosinophil percentage decreased, Glomerular filtration rate, Haematocrit normal, Haemoglobin normal, Hypoaesthesia, Immature granulocyte count increased, Immature granulocyte percentage increased, Injection site hypoaesthesia, Injection site pain, Lymphocyte count decreased, Lymphocyte percentage decreased, Mean cell haemoglobin concentration normal, Mean cell haemoglobin normal, Mean cell volume normal, Mean platelet volume normal, Monocyte count normal, Monocyte percentage decreased, Nausea, Neutrophil count increased, Neutrophil percentage increased, Pain in extremity, Platelet count normal, Protein total normal, Red blood cell count normal, Red blood cell nucleated morphology, Red cell distribution width increased, Throat tightness, Troponin normal, White blood cell count normal
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (broad), Haematopoietic leukopenia (narrow), Peripheral neuropathy (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad), Hypersensitivity (broad), Myelodysplastic syndrome (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypokalaemia (narrow), Sexual dysfunction (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes

Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Amitriptyline 25 mg PO QHS Bupropion XL 150 mg PO Daily Cyanocobalamin 1000 mcg PO once weekly Ferrous sulfate DR 324 mg PO Daily Levothyroxine 100 mcg PO Daily Methocarbamol 500 mg PO TID prn muscle spasms Scopolamine 1.5 mg patch transder
Current Illness:
Preexisting Conditions: Anemia Asthma Depression Hypothyroidism secondary to thyroid cancer s/p thyroidectomy Hypertension Migraines Vertigo
Allergies: No known allergies
Diagnostic Lab Data: Chest X-ray (12/30/20 @ 1737): The cardiac and mediastinal silhouette are unremarkable. The lungs are free of infiltrates and masses. I see no acute chest disease on today"s film. CBC w/diff (12/30/2020 @ 1658) WBC: 9.0 RBC: 4.38 Hemoglobin: 13.3 Hematocrit: 39.8 MCV: 90.9 MCH: 30.4 MCHC: 33.4 RDW SD: 40.4 Platelet: 264 MPV: 9 NRBC Percent: 0.0 NRBC Absolute: 0.000 Absolute Neutrophil: 8.4 (H) Lymphocyte Absolute: 0.4 (L) Monocytes Absolute, Automated: 0.11 Absolute Eosinophil: 0.00 (L) Basophil Absolute: 0.03 Absolute Immature Granulocytes: 0.03 (H) Neutrophil percent: 93.6 (H) Lymphocyte percent: 4.6 (L) Monocyte percent: 1.2 Eosinophil percent: 0.0 Basophil percent: 0.3 Immature Granulocytes percent: 0.30 (H) CMP (12/30/2020 @ 1658) Sodium: 136 Potassium: 3.3 (L) Chloride: 106 CO2: 21 Anion gap: 9 Glucose: 137 (H) BUN: 15 Creatinine: 1.01 BUN/Creatinine Ratio: 14.9 Calcium: 9.2 Protein total: 7.9 Albumin: 4.6 AST: 12 ALT: 17 ALP: 50 eGFR non-African American: $g60.0 eGFR African American: $g60.0 Bilirubin total: 0.4 Magnesium: 2.0 TSH: 2.52 Cardiovascular (12/30/2020 @ 1658) Troponin: <0.015
CDC Split Type:

Write-up: Presented to the ED after developing chest tightness, cough, lightheadedness, and throat closing sensation. She received the Moderna COVID-19 vaccine on the morning of presentation. Within 15 minutes of receiving the vaccine she developed pain and numbness, starting at the injection site traveling down the ulnar aspect of her arm, and nausea. Over the next several hours she continued to develop worsening nausea, chest tightness, cough, lightheadedness, and the sensation that her throat closing. She took PO Benadryl 25mg; however, her symptoms were not alleviated. She was subsequently evaluated in the ED. ? Received PO Benadryl 25mg, IV Benadryl 25mg, Epinephrine 0.3mg x 2, IV Famotidine 20mg, IV Solumedrol 125mg & 60mg, DuoNebs x 3, Racepinephrine x 1.

VAERS ID: 919620 (history)
Form: Version 2.0
Age: 81.0
Sex: Male
Location: Washington
Vaccinated: 2020-12-01
Onset: 2021-01-01
Days after vaccination: 31
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot /	Site /

	Dose	Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM
UNK: VACCINE NOT SPECIFIED (NO BRAND NAME) / UNKNOWN MANUFACTURER	EL0140 / 1	UN / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Body temperature increased, Cardiac failure, Pyrexia
SMQs:, Cardiac failure (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Mucus Relief Tablet 400 MG (GuaiFENesin) Give 1 tablet by mouth every 8 hours as needed for congestion Prescriber Entered Active01/02/202101/04/2021 Omeprazole Capsule Delayed Release 40 MG Give 1 capsule by mouth in the morning for GERD Ph
Current Illness: N/A
Preexisting Conditions: Dementia, Urinary Incontinence, depression, Edema
Allergies: No
Diagnostic Lab Data:
CDC Split Type:

Write-up: Decompensation and temp 103.6.

VAERS ID: 919624 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Asthenia, Blood pressure decreased, Chills, Decreased appetite, Dizziness, Fatigue, Headache, Hot flush, Lymphadenopathy, Mobility decreased, Nausea, Pain in

extremity, Paraesthesia, Speech disorder, Syncope
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Acute pancreatitis (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Parkinson-like events (broad), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: no
Current Illness: no
Preexisting Conditions: no
Allergies: Penicillin
Diagnostic Lab Data: no
CDC Split Type:

Write-up: Around 10 or 11 pm, arm pain, chills, fatigue, headache, nausea, swollen lymph nodes, lightheadedness, fainted in tub. Next day, fatigue all day, couldn''t talk, or eat. Went back to bed again. The following day hot flashes, weakness. went to ER. Felt like blood pressure was dropping, tingling in legs, difficulty lifting her head.

VAERS ID: 919633 (history)
Form: Version 2.0
Age: 49.0
Sex: Female
Location: Mississippi
Vaccinated: 2020-01-04
Onset: 2020-12-30
Days after vaccination: 361
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Alanine aminotransferase normal, Angiogram cerebral normal, Anion gap decreased, Arteriogram carotid normal, Aspartate aminotransferase normal, Basophil count

normal, Basophil percentage, Blood albumin normal, Blood alkaline phosphatase normal, Blood bilirubin normal, Blood calcium normal, Blood chloride normal, Blood creatinine normal, Blood glucose normal, Blood potassium normal, Blood sodium normal, Blood urea nitrogen/creatinine ratio, Blood urea normal, Carbon dioxide normal, Chest X-ray normal, Computerised tomogram head normal, Electrocardiogram, Eosinophil count normal, Eosinophil percentage, Facial paralysis, Facial paresis, Feeling hot, Glomerular filtration rate, Haematocrit normal, Haemoglobin normal, Hypoaesthesia, Immature granulocyte count increased, Immature granulocyte percentage increased, Immediate post-injection reaction, Injection site hypoaesthesia, Injection site warmth, Lymphocyte count normal, Lymphocyte percentage decreased, Mean cell haemoglobin concentration normal, Mean cell haemoglobin decreased, Mean cell volume normal, Mean platelet volume increased, Monocyte count normal, Monocyte percentage, Muscular weakness, Nausea, Neutrophil count normal, Neutrophil percentage increased, Palpitations, Platelet count normal, Protein total normal, Red blood cell count normal, Red blood cell nucleated morphology, Red cell distribution width, White blood cell count normal
SMQs:, Rhabdomyolysis/myopathy (broad), Acute pancreatitis (broad), Haematopoietic leukopenia (broad), Peripheral neuropathy (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hearing impairment (broad), Hypersensitivity (narrow), Myelodysplastic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Albuterol 90 mcg/actuation 2 puffs inhaled q4h prn sob/wheezing Aspirin EC 81 mg PO Daily Bupropion SR 150 PO Daily Calcium carbonate + Vitamin D 600mg PO Daily Cinnamon Bark 2 capsules PO Daily prn Clonazepam 0.5 mg PO Daily prn anxiety Cl
Current Illness: N/A
Preexisting Conditions: Asthma Fibrocystic breast disease Hyperlipidemia Irritable bowel syndrome Insomnia Migraines Polycystic ovarian syndrome
Allergies: Latex gloves - Anaphylaxis Iodine - Swelling Shellfish - Unknown Oseltamivir - Unknown Tramadol - Itching, N/V
Diagnostic Lab Data: CT Head w/o contrast (12/31/20 @ 1813) READING: I see no evidence of hemorrhage, mass effect, abnormal attenuation, extra-axial lesions, ventriculomegaly or other suspicious intracranial pathology. Included paranasal sinuses, mastoids and middle ears are clear. Calvarium intact. IMPRESSION: no acute intracranial pathology Chest X-Ray (12/31/20 @ 1821) READING: No suspicious cardiomediastinal pathology. No infiltrates or other suspicious pulmonary or pleural pathology. I see no acute abnormality. CT Angiogram w/o contrast (12/31/20 @ 1816) READING: Both common carotid arteries and the vertebral arteries are patent and normal. Do not see any plaque or stenosis. Both carotid bifurcations are normal. The internal carotid arteries remain patent throughout the course without stenosis or occlusion in the basal artery is normal. Anterior posterior cerebral arteries widely patent without stenosis or occlusion. No arterial venous malformation or

aneurysm or other vascular abnormalities. IMPRESSION: No arterial stenotic or occlusive disease of the head or neck EKC 12-Lead (12/31/20 @ 1728) Systolic Blood Pressure: 172 Diastolic Blood Pressure: 96 Vent Rate: 93 Atrial Rate: 93 PR Interval: 150 QRS Interval: 78 QT Interval: 356 QTC Interval: 442 P Axis: 66 R Axis: 25 T Axis: 22 CBC w/diff (12/31/20 @ 1845) WBC: 6.4 RBC: 5.28 Hemoglobin: 14.2 Hematocrit: 43.7 MCV: 82.8 MCH: 26.9 (L) MCHC: 32.5 RDW SD: 37.8 Platelet: 197 MPV: 11 (H) NRBC Percent: 0.0 NRBC Absolute: 0.000 Absolute Neutrophil: 4.9 Lymphocyte Absolute: 1.1 Monocytes Absolute, Automated: 0.38 Absolute Eosinophil: 0.07 Basophil Absolute: 0.05 Absolute Immature Granulocytes: 0.01 (H) Neutrophil percent: 75.2 (H) Lymphocyte percent: 16.8 (L) Monocyte percent: 5.9 Eosinophil percent: 1.1 Basophil percent: 0.8 Immature Granulocytes percent: 0.20 (H) CMP (12/31/20 @ 1845) Sodium: 138 Potassium: 3.9 Chloride: 105 CO2: 28 Anion gap: 5 (L) Glucose: 82 BUN: 19 (H) Creatinine: 0.88 BUN/Creatinine Ratio: 21.6 Calcium: 9.8 Protein total: 7.8 Albumin: 3.9 AST: 24 ALT: 23 ALP: 71 eGFR non-African American: $g60.0 eGFR African American: $g60.0 Bilirubin total: 0.8
CDC Split Type:

Write-up: Presented to the ED with cc of left sided facial and LUE numbness and weakness x 1 days. Patient received her COVID-19 vaccination on 12/30/2020 around 1PM. Immediately after the injection in her left shoulder, she began to feel warmth and numbness in her left shoulder, arm, neck, face, and chest. She reports later experiencing nausea, palpitations, and left arm weakness. Her symptoms persisted, and her family noted a left sided facial droop which prompted her ED visit. In the ED, patient was noted to have some left sided facial droop and left arm and leg weakness. CT head and CTA showed no acute abnormalities. Tele-neurology was consulted who recommended admission to rule out acute stroke. Ultimately, work up was negative and symptoms resolved. Symptoms appear to be related to the vaccine.

VAERS ID: 919665 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: New Jersey
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Arthralgia, Blood test, Chest X-ray, Nausea, Pyrexia
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Arthritis (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Synthroid Claritin Dexilent Terbinafine Vitamin D & B12
Current Illness:
Preexisting Conditions: Hypothyroid GERD
Allergies: Bactrim
Diagnostic Lab Data: Blood work 12/30/20 Chest X-ray 12/30/20
CDC Split Type:

Write-up: Severe joint aches, fever-type symptoms, nausea

VAERS ID: 918705 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-12
Onset: 2020-12-12
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Cerebellar infarction, Confusional state, Disorientation, Dizziness, Facial paralysis, Ischaemic stroke, Magnetic resonance imaging
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Ischaemic central nervous system vascular conditions (narrow), Dementia (broad), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hearing impairment (broad), Vestibular disorders (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days

Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Hypertension; Small vessel disease of diabetes mellitus
Allergies:
Diagnostic Lab Data: Test Name: MRI; Result Unstructured Data: Test Result:showed right cerebellar infarct
CDC Split Type: GBPFIZER INC2020517517

Write-up: Cerebellar infarction; Confusion; Facial droop; Ischaemic stroke; Dizziness; Disorientated; This is a spontaneous report from a contactable pharmacist downloaded from the Medicines Agency (MA) Regulatory Authority-WEB GB-MHRA-EYC 00235714, Safety Report Unique Identifier GB-MHRA-ADR 24544651. A 58-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), intramuscular on 12Dec2020 at single dose (reported as 30ug) for COVID-19 vaccination. Medical history included hypertension and small vessel disease of diabetes mellitus unknown if ongoing. Concomitant medication included amlodipine. The patient experienced dizziness on 12Dec2020, disorientated on 12Dec2020, cerebellar infarction on unknown date, confusion on unknown date, ischaemic stroke on 12Dec2020, facial droop on an unspecified date. The events were all serious as causing hospitalization from 17Dec2020, medically significant, life threatening. Signs/Symptoms: Right eye drooping; slight confusion; right arm reflex slightly impaired. The patient underwent lab tests and procedures which included magnetic resonance imaging: showed right cerebellar infarct on unknown date. The outcome of event ischaemic stroke was recovering while for other events was unknown. No follow-up attempts are possible, information on batch number cannot be obtained.

VAERS ID: 918706 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-19
Onset: 2020-12-01
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Angiogram, Atrial fibrillation, Blood pressure measurement, Blood test, Cardiac arrest, Cold sweat, Computerised tomogram, Dyspnoea, Investigation, Malaise, Oxygen saturation, Pallor, Respiration abnormal, Unresponsive to stimuli
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension

(broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Respiratory failure (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; CO-CODAMOL
Current Illness: Smoker
Preexisting Conditions: Medical History/Concurrent Conditions: Aortic valve sclerosis; Cardiovascular collapse; Mitral regurgitation; Ventricular tachycardia
Allergies:
Diagnostic Lab Data: Test Date: 202012; Test Name: Angiogram; Result Unstructured Data: Test Result:CT angiogram showed rib fractures, no PE.; Comments: CT angiogram showed rib fractures, no PE.; Test Date: 20201220; Test Name: Blood pressure; Result Unstructured Data: Test Result:Good; Test Date: 202012; Test Name: Blood test; Result Unstructured Data: Test Result:Normal; Test Date: 202012; Test Name: CT scan; Result Unstructured Data: Test Result:Emphysema; Test Date: 202012; Test Name: Investigation; Result Unstructured Data: Test Result:New onset atrial fibrillation; Comments: New onset atrial fibrillation; Test Date: 20201220; Test Name: Oxygen saturation; Test Result: 97 %
CDC Split Type: GBPFIZER INC2020518031

Write-up: Cardiac arrest; Unresponsive to stimuli; Clammy; Gasping; Pale; look unwell; agonal breathing; atrial fibrillation; This is a spontaneous report from a contactable physician downloaded from the Regulatory Authority - GB-MHRA-EYC 00235831, Safety Report Unique Identifier GB-MHRA-ADR 24545081. An 85-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, batch/ lot no: EJ0553), intramuscularly on 19Dec2020 10:37 at 0.3 mL, single for COVID-19 vaccination. Medical history included Cardiovascular collapse from 2016 and not ongoing, ongoing Smoker, Ventricular tachycardia from 2016 and not ongoing, Aortic valve sclerosis from an unknown date and unknown if ongoing, Mitral regurgitation from an unknown date and unknown if ongoing (past history of cardiac disease, more recently aortic valve sclerosis further complicated with mitral regurgitation). Concomitant medication included bisoprolol, tamsulosin, omeprazole, codeine phosphate, paracetamol (CO-CODAMOL). The patient experienced cardiac arrest with outcome of recovered on 19Dec2020, unresponsive to stimuli with outcome of recovered in Dec2020, clammy with outcome of recovered in Dec2020, gasping with outcome of recovered in Dec2020, pale with outcome of recovered in Dec2020, look unwell with outcome recovered in Dec2020, agonal breathing with outcome recovered in Dec2020, all on 19Dec2020, broken ribs in Dec2020 with outcome unknown, atrial fibrillation in Dec2020 with outcome unknown. The patient had vaccine at 10:37 AM, sat in waiting room and noted to look unwell at 10:50. Doctor was called by other patients as subject seemed unwell, gasping, looked pale, became clammy, unresponsive with agonal breathing. Agonal gasps so cardiopulmonary resuscitation (CPR) commenced. He had cardiac arrest and received CPR and defibrillation. He had non-shockable rhythm. He was cannulated, given oxygen and adrenaline, and 5 cycles CPR, then became responsive. He was placed in the recovery positron, and given high flow oxygen until

the ambulance arrived. The patient had some chest pain overnight, consistent with CPR he received. Computerised tomography (CT) angiogram showed rib fractures and no pulmonary embolism. The patient has new onset atrial fibrillation. No features of anaphylaxis at all but treated with anaphylaxis kit anyway. Return of spontaneous circulation and now in hospital. Has history of previous episodes of collapse and ventricular tachycardia. Post vaccine, treating physician confirmed no allergic type reaction but patient was also treated as if he had anaphylaxis. Ambulance was called and by the time the ambulance arrived he recovered, orientated and able to talk. He was hospitalised for observation and has broken ribs due to CPR. It was agreed that the vent of cardiac arrest observed within 15 minutes after vaccination was coincidental. Time of onset is short, adverse event could potentially be associated with severe allergic reaction, however despite being treated for anaphylaxis, no other symptoms of anaphylaxis/severe allergic reactions were reported. Patient had several concomitant risk factors for cardiac arrest including age, smoking and past history of cardiac disease. No signs of anaphylaxis in the events, the patient has collapsed previously, his daughter says this happened at a funeral, and considers it related to emotions. CT scan shows emphysema. Nobody at health centre thought this event was related to the vaccination. Blood results look good in hospital, "pretty much normal". Patient"s oxygen saturations were 97% at 09:30 20Dec2020, blood pressure was good and was awake. The patient underwent lab tests and procedures which included angiogram: CT angiogram showed rib fractures, no PE in Dec2020; blood pressure: good on 20Dec2020; blood test: normal in Dec2020; CT (computerised tomogram) scan: emphysema in Dec2020; investigation: new onset atrial fibrillation in Dec2020; oxygen saturation: 97 % on 20Dec2020. All events were serious per hospitalization, life threatening. No follow-up attempts possible. No further information expected.

VAERS ID: 918714 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-15
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Computerised tomogram head, Computerised tomogram thorax, Lip swelling, Pyrexia, SARS-CoV-2 test, Swollen tongue, Vomiting
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? Yes

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; ; ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Acute kidney injury; Anaemia; Anxiety; Clostridium difficile infection; Depression; Fracture of ankle (left ankle); Hypertension; Infection MRSA; Learning disorder (learning difficulties); Multinodular goitre; Osteoporosis; Stroke
Allergies:
Diagnostic Lab Data: Test Name: Brain CT; Result Unstructured Data: Test Result:unknown results; Test Name: Chest CT; Result Unstructured Data: Test Result:unknown results; Test Date: 20201215; Test Name: COVID-19 virus test; Test Result: Negative ; Comments: No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2020517590

Write-up: Lip swelling; Vomited; Pyrexia; Tongue swelling non-specific; This is a spontaneous report from a contactable other health professional downloaded from the Medicines Agency (MA) Regulatory Authority-WEB. This is a report received from the regulatory authority. The regulatory authority report number is GB-MHRA-WEBCOVID-20201217093758; other case identifier number: GB-MHRA-ADR 24544180. A 75-year-old female patient received the bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on 15Dec2020 at 14:00 at a single dose for COVID-19 immunization. Medical history included Stroke, anxiety, depression, multinodular goitre, infection methicillin-resistant Staphylococcus aureus (MRSA), Fracture of ankle (left ankle), anaemia, Clostridium difficile infection, acute kidney injury, hypertension, osteoporosis, learning disorder (learning difficulties); all from an unknown date and unknown if ongoing. Concomitant medications included thiamine (MANUFACTURER UNKNOWN), ferrous fumarate (MANUFACTURER UNKNOWN), simvastatin (MANUFACTURER UNKNOWN), colecalciferol (MANUFACTURER UNKNOWN), bisoprolol (MANUFACTURER UNKNOWN) , carbimazole (MANUFACTURER UNKNOWN), amlodipine (MANUFACTURER UNKNOWN), clopidogrel (MANUFACTURER UNKNOWN), paracetamol (MANUFACTURER UNKNOWN); all taken for an unspecified indication from an unspecified date to an unspecified date. The patient experienced the following events and outcomes: lip swelling (hospitalization, life threatening) on 15Dec2020 with outcome of recovered on an unspecified date, vomited (hospitalization, life threatening) on 15Dec2020 with outcome of recovered on an unspecified date, pyrexia (hospitalization, life threatening) on 15Dec2020 with outcome of recovering, tongue swelling non-specific (hospitalization, life threatening) on 15Dec2020 with outcome of recovered on 17Dec2020. The events developed 40 minutes after the vaccination. The patient was hospitalized on unspecified dates. The patient had not tested positive for COVID-19 since having the vaccine. The patient underwent lab tests and procedures which included Brain computerised tomogram (CT): unknown results on an unspecified date, Chest CT: unknown results on an unspecified date, COVID-19 virus test: negative on 15Dec2020 No - Negative COVID-19 test. Additional information: "Unsure if patient has had symptoms associated with COVID-19. Patient is not enrolled in clinical trial." The batch/lot numbers for the vaccine, PFIZER-BIONTECH COVID-19 MRNA VACCINE, were not provided and will be requested during

follow up.

VAERS ID: 918718 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6796 / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Defaecation urgency, Flatulence, Flushing, Gastrointestinal hypermotility, Mucosal dryness, Tachycardia
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Gastrointestinal nonspecific dysfunction (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (narrow), Noninfectious diarrhoea (narrow), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GRPFIZER INC2020516043

Write-up: moderate anaphylactic reaction; flushing; tachycardia; mucosal dryness; hypermotility of the intestine (gases); hypermotility of the intestine (gases); urgency to defecate; This is a spontaneous report from two contactable physicians (initially from an intern (patient) reported at 28Dec2020, at 19:00 local time and then from the vaccination coordinator of the hospital). An approximately 30-year-old female patient (HCP, and intern herself) was administered the first dose of bnt162b2 (COMIRNATY, Solution for injection; lot number: EJ6796, expiration date: 30Apr2021), intramuscularly (at the deltoid) on 28Dec2020 at single

dose for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. On 28Dec2020, after vaccination, the patient developed tachycardia, flushing, mucosal dryness, intestinal hypermotility (initially gases and then urgency to defecate) categorized as moderate anaphylactic reaction (anaphylactic reaction). The patient initially was administered cortisone NOS that didn"t help and then she was administered adrenaline. She recovered quickly after the adrenaline administration. The patient required hospitalization for 5 - 6 hours, which was prophylactically extended to overnight hospitalization as a precaution. The vaccination center was at a hospital. The patient was not admitted to the Intensive Care Unit. The organ systems affected were: Cardiovascular, Dermatological/Mucosal, Gastrointestinal. Respiratory system was not affected. It was not known whether the patient have a history of any previous allergies to specific products or any conditions indicative of an allergy. Past product history was not known. It was unknown whether the patient received any recent vaccines for any other conditions prior to the event being reported. The same event was reported earlier on live TV (28Dec2020 at 18:15 local time), during the Ministry of Health established regular Press Conference on the pandemic update, where one of the speakers (general secretary of primary health treatment) mentioned that: "Out of the 471 HCPs vaccinated by 1600 h today, only one case was reported of non-serious allergic reaction that was treated successfully". Later, at the news it was reported by journalists that the patient was a physician who was intern at a COVID-19 reference hospital (the name of the hospital was provided) and she had a mild allergic reaction that did not require hospitalization (as reported). It was also reported that the hospital also filed a yellow card to the local HAs. The clinical outcome of the events was recovered on 28Dec2020. Follow-up activities not initiated. No consent for further follow-up obtained.; Sender"s Comments: The information available in this report is limited and does not allow a medically meaningful assessment of the case. In particular the following relevant information is not available: complete medical history and complete demographics, concomitant medications and full diagnostic workup. There is a plausible time relationship between vaccine administration and onset of events that are only partially suggestive of an anaphylactic reaction. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 919827 (history)
Form: Version 2.0
Age: 45.0
Sex: Male
Location: California
Vaccinated: 2020-12-22
Onset: 2020-12-23
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	11J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain lower, Abdominal pain upper, Appendicectomy, Appendicitis, Computerised tomogram abdomen abnormal, Discomfort, Nausea, Pain
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal

nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none
Preexisting Conditions: none
Allergies: penicillin
Diagnostic Lab Data: evaluated by urgent care physicians and determined to likely be presenting with an appendicitis. Diagnosis of acute appendicitis confirmed and surgery performed to remove appendix.
CDC Split Type:

Write-up: Began experiencing nausea and general stomach pains the morning after receiving the vaccine. After one day of pain and discomfort I woke the following morning (~44hrs after receiving the vaccine) to extreme acute abdominal pain in the lower right abdomen. Went to Urgent Care facility and was diagnosed by CT scan as having acute appendicitis. An emergency appendectomy was scheduled and performed for later that evening. I stayed at the Hospital overnight on 12/24/2020 post operatively on IV antibiotics to recover from the appendectomy. Was discharged from the hospital on 12/25/2020 and have been recovering for about 10 days now with limitted activity.

VAERS ID: 920224 (history)
Form: Version 2.0
Age: 35.0
Sex: Male
Location: Texas
Vaccinated: 2020-12-18
Onset: 2020-12-01
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Chills, Computerised tomogram abnormal, Cough, Dyspnoea, Hypoxia, Intensive care, Lung infiltration, Myalgia, Pain, Pyrexia, SARS-CoV-2 test positive
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Interstitial lung disease (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression

(broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Date: 202012; Test Name: CT scan; Result Unstructured Data: Test Result:showed extensive infiltration in the lungs; Test Date: 20201218; Test Name: O2 saturation; Result Unstructured Data: Test Result:80%; Test Date: 20201226; Test Name: COVID test; Test Result: Positive
CDC Split Type: USPFIZER INC2020518143

Write-up: had a positive COVID test; had a positive COVID test; O2 Saturation of 80% / Hypoxia; shortness of breath; He has a CT scan which showed extensive infiltration in the lungs; muscle pain; chills; body aches; low grade fever; cough; This is a spontaneous report from a contactable physician (pulmonary medicine). This physician reported similar events for 2 patients. This is 1st of 2 reports. A 35-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 18Dec2020 at single dose for covid-19 immunization. There were no medical history and concomitant medications. Caller stated that his close friend who was ER physician (front line worker) and within 24 hours after receiving the COVID vaccine, developed COVID or symptoms of COVID. Patient received the COVID vaccine on 18Dec2020 and the same night patient started with a low grade fever, body aches, chills, muscle pain, shortness of breath, cough, O2 saturation of 80% (hypoxia) and was in the intensive care unit now. Patient swore this was related to the vaccine. This patient tested positive for COVID. He had a CT (computerised tomogram) scan which showed extensive infiltration in the lungs in Dec2020. Patient was admitted to the hospital on 24Dec2020 and then was moved to the ICU 2 days later, on 26Dec2020. Caller thought patient had a positive COVID test at another hospital. Caller did know that tested positive at the current hospital on 26Dec2020 which was done to confirm the previous positive test. Caller thought patient had his first positive COVID test either the same day or the next day after receiving the vaccine. Event of O2 Saturation of 80% / hypoxia was reported as hospitalization from 24Dec2020 and life threatening; infiltration in the lungs and shortness of breath caused hospitalization from 24Dec2020, muscle pain, chills and positive COVID test was reported as medically significant; and other events were reported as non-serious. Outcome of O2 saturation of 80% / hypoxia and shortness of breath was not recovered, outcome of cough was recovering; and outcome of other events were unknown. Information about lot/batch number has been requested. ; Sender"s Comments: Based on the information currently available, a lack of efficacy with suspected vaccine BNT162B2 in this patient cannot be completely excluded.,Linked Report(s) : US-PFIZER INC-2020519020 same

reporter/drug , different patient/AE.

VAERS ID: 920225 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: California
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / OT

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Viral test negative
SMQs:, Anaphylactic reaction (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: VITAMIN C [ASCORBIC ACID]; VITAMIN D [COLECALCIFEROL]
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Date: 20201218; Test Name: Nasal Swab; Test Result: Negative
CDC Split Type: USPFIZER INC2020518186

Write-up: Anaphylactic reaction requiring two doses of Epinephren to control. Still having issues; other vaccine same date product received; This is a spontaneous report from a contactable nurse (patient). A 54-year-old female patient received her first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), intramuscularly on right arm at 12:30 PM on 18Dec2020 at single dose for COVID-19 immunization, and first dose of other Pfizer vaccine same date product (other vaccine same date lot number: elt9899) on right deltoid on 18Dec2020 for unknown indication. Medical history reported as none. Concomitant medication included vitamin C and colecalciferol (VITAMIN D). The patient experienced anaphylactic reaction at 12:30 PM on 18Dec2020 requiring two doses of epinephren to control. still having issues, resulted in: Doctor or other healthcare professional office/clinic

visit, Hospitalization in Dec2020. days hospitalization: 1. The patient received treatment: 2 doses of epinephrine, solumedrol, benadryl, IV and O2 for event anaphylactic reaction to vaccine. The outcome of anaphylactic reaction was not recovered. Lot/Batch and Expiration date has been requested.; Sender"s Comments: The information available in this report is limited and anecdotal and does not allow a medically meaningful assessment of the case. There is a plausible temporal association between vaccines administration and onset of the reported event. It is unclear what is the nature of the vaccine co-administered with BNT162b2. Currently no information is available on the co-administration of the Pfizer-BioNTech COVID-19 Vaccine with other vaccines.

VAERS ID: 920280 (history)
Form: Version 2.0
Age: 54.0
Sex: Male
Location: Unknown
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Military **Purchased by:** ?
Symptoms: Asthenia, Dizziness, Hyperhidrosis, Hypertension
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Hypertension (narrow), Vestibular disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Multivitamins
Current Illness: DENIES
Preexisting Conditions: DENIES
Allergies: DENIES
Diagnostic Lab Data:
CDC Split Type:

Write-up: 54yoM presented for a sudden-onset diaphoresis and loss of body strength. Pt had received COVID-19 vaccine this morning at the clinic and noted severe sweating all over the body while lying in bed @ approx 2030. Then pt described severe loss of body strength with dizziness as he was trying to sit up and get to his feet from the floor. Denied palpitation, chest

pain, headache, N/V, LOC. His symptoms continued for 20mins until he drank a cup of water. At approx 2050, smoked E-cigarette and arrived at facility at 2100. On arrival, pt denied any symptoms, however, vitals indicated severe hypertension: BP 188/99, HR 56, T 97.6 F. Pt given 81mg ASA x2 and immediately transported to hospital.

VAERS ID: 920317 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: New York
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Heart rate increased, Pyrexia, Rhonchi, Sputum abnormal
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: buspirone, prednisone, Vit D3, Vit B1, Multivitamin, Ensure Plus, Fentanyl Patch, Levothyroxine, Metoprolol, Ipratropium-Albuterol SVN
Current Illness: Sepsis, Pneumonia, UTI, Respiratory Failure, COPD
Preexisting Conditions: CA lymph nodes, liver, brain lung and base of tongue, COPD, CKD, anemia, DM Type 2, Vit D3 Deficiency
Allergies: Bactrim DS, bupropion (bulk), fluoxetine, piperacillan-tazobactam, remeron, latez
Diagnostic Lab Data: none
CDC Split Type:

Write-up: Rhonchi, frothy sputum, low grade temp, elevated HR, 12/30 MD assessed 1:00PM and increased prednisone and added Cefdinir. Sent to hospital 12/30 at approximately 6:30PM with worsening symptoms

VAERS ID: 920388 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-16
Onset: 2020-12-16
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Chills, Diarrhoea, Headache, Meningitis viral, Nausea, Photophobia, Pyrexia, Transaminases increased, Vomiting
SMQs:, Liver related investigations, signs and symptoms (narrow), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Noninfectious meningitis (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Glaucoma (broad), Corneal disorders (broad), Retinal disorders (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: craniotomy in 1998 due to meningioma
Allergies: codeine
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient received her Vaccine on 12/16/2020. Afterwards she developed symptoms of fever, chills, diarrhea, nausea, vomiting and headache that became worse over time and on day os presentation to our hospital on 1/2/2021 she was having photophobia. Current headache at time of admission had been persistent for over a week. Patient has no immunocompromising risk factors and was diagnosed with confirmed CMV meningitis. She was also admitted with transaminitis.

VAERS ID: 920628 (history)
Form: Version 2.0
Age: 63.0
Sex: Female
Location: New York
Vaccinated: 2021-01-02
Onset: 2021-01-02
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0142 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Chills, Colostomy, Computerised tomogram abdomen, Computerised tomogram abnormal, Intestinal resection, Large intestine perforation, Pain in extremity, Pyrexia
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Malignancy related therapeutic and diagnostic procedures (narrow), Gastrointestinal perforation, ulcer, haemorrhage, obstruction non-specific findings/procedures (broad), Gastrointestinal perforation (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Ischaemic colitis (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Tylenol 350 mg as needed Valium 5 mg as needed
Current Illness:
Preexisting Conditions: pancreatic cysts colonic polyps last colonoscopy 9/2020 and one polyp removed ATM gene mutation hyperlipidemia DCIS of breast in 1/2015 s/p lumpectomy and xrt thyroid nodules vertigo
Allergies: Compazine led to restless leg syndrome Penicillin anaphylaxis Sulfamethoxazole rash
Diagnostic Lab Data: CT scan of abdomen and pelvis 1/3/2021
CDC Split Type:

Write-up: 6-7 hours after the vaccine she developed arm pain, fever and chills. About an hour later she started to have abdominal pain which worsened over the course of the day to excruciating. She went to the Emergency Room where a CT scan revealed a perforation of

her sigmoid colon and had a resection of the area of the colon and a diverting colostomy surgery done the evening of 1/3/2021.

VAERS ID: 920719 (history)
Form: Version 2.0
Age: 36.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-23
Onset: 2020-12-28
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Blood test, Epistaxis, Immune thrombocytopenia, Lethargy, Menorrhagia, Oral blood blister, Petechiae, Platelet count decreased
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none
Preexisting Conditions: none
Allergies: none
Diagnostic Lab Data: blood work
CDC Split Type:

Write-up: starting to feel lethargic and weak. Had menses with increased blessed. Called physician to have blood work done to see if I was experiencing anemia. Blood work complete on 12/31/2020. On 1/3/2021, I woke up with blood blisters all over the inside of my mouth and petechia on my trunk and bilateral upper and lower extremities. I called my primary physician to report the symptoms. He suggested to go to the ER if my symptoms worsened. Later that

evening I started with a nose bleed and did go to the ER. Upon arrival to the ER, my platelet count was 9. I was admitted to the hospital and diagnosed with ITP.

VAERS ID: 920726 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: Kentucky
Vaccinated: 2020-12-29
Onset: 2020-12-31
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dysarthria, Electrocardiogram normal, Full blood count normal, Hypoaesthesia, Injection site rash, Lymphadenopathy, Magnetic resonance imaging brain normal, Magnetic resonance imaging neck, Platelet count normal, Troponin normal
SMQs:, Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: 12/31/20 MRI of brain, Carotid Artery CBC, Platelet panel, Troponin, EKG
CDC Split Type:

Write-up: 12/31/20 around 11am Numbness in right hand and right cheek, 5 minutes later, slurred speech. Episode lasted approximately 10 minutes. Treated in ER. Labs, MRI, all normal 1/3/21 Swollen lymph node to left axilla 1/4/21 Rash to injection site

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VAERS ID: 920771 (history)

Form: Version 2.0
Age: 94.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Tachycardia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: NKA
Diagnostic Lab Data:
CDC Split Type:

Write-up: Tachycardia, resident was sent out to the hospital for evaluation on 12/30/2020 and came back to the facility on 12/31/2020.

VAERS ID: 920784 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: Indiana
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot /	Site / Route

	Dose	
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	LA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Anxiety, Blood test, Burning sensation, Central nervous system lesion, Chest X-ray, Differential white blood cell count, Dizziness, Dysphagia, Ear pain, Electrocardiogram, Erythema, Flushing, Full blood count, Headache, Hypertension, Hypoaesthesia, Hypoaesthesia oral, Magnetic resonance imaging brain abnormal, Metabolic function test, Mobility decreased, Palpitations, Paraesthesia, Paraesthesia oral, Peripheral swelling, Pharyngeal swelling, Swelling face
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Angioedema (narrow), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypertension (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Sexual dysfunction (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: 12/23/20 EKG, X-ray of lungs, Lab Blood work 1/1/2021 Basic Metabolic Panel, blood count with auto diff, ECG 12- Lead, MRI Brain WO contrast
CDC Split Type:

Write-up: Anaphylactic Reaction, facial swelling, facial Redness, Face felt like it was burning, face flushing, throat swelling, heart palpitations, trouble swallowing , feet swelling, light headed, anxiety. Hospitalized from the 12/23/20 to 12/26/2020 . Medications now on Epinephrine, diphenhydramine, cetirizine, famotidine, prednisone, lorazepam, cephalexin. on 1/1/2021 was taken to E.R. by ambulance around 11:00 am left hand was tingle started to go numb traveled up my arm into left side of my face ,ear, tongue, and then down to the left side of my leg and into left foot, could not move left side of body for a good 7 to 8 mins then went away transferred to ambulance enroute to ER blood pressure was high and and started having right ear pain and right side frontal severe headache, arrived to ER and was given diphenhydramine ,ketorolac, metoclopramide HCl, lorazepam. MRI was ordered and Neurologist found two small lesions on right side of frontal brain, following up now with

neurologist. added more meds naproxen

VAERS ID: 920805 (history)
Form: Version 2.0
Age: 45.0
Sex: Male
Location: Ohio
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Acute kidney injury, Alanine aminotransferase increased, Aspartate aminotransferase increased, Blood alkaline phosphatase increased, Blood creatine increased, Blood pressure increased, Chest discomfort, Head discomfort, Liver function test increased, Palpitations, Throat clearing, Throat irritation, Vision blurred
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Liver related investigations, signs and symptoms (narrow), Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Biliary system related investigations, signs and symptoms (broad), Glaucoma (broad), Hypertension (narrow), Cardiomyopathy (broad), Lens disorders (broad), Retinal disorders (broad), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: NKDA
Diagnostic Lab Data: AP 144, AST 51, ALT 178 Creatinine 1.41

CDC Split Type:

Write-up: palpitations, chest tightness/heaviness, scratchy throat/frequent throat clearing, head heaviness, blurred vision, elevated blood pressure. Evaluated in ED received solumedrol, benadryl. Discharged home and returned to hospital as direct admit due to continued symptoms. Pertinent labs reveled AKI, elevated LFTs. AKI and LFTs improved with IVFs.

VAERS ID: 920879 (history)
Form: Version 2.0
Age: 36.0
Sex: Female
Location: Wisconsin
Vaccinated: 2020-12-23
Onset: 2020-12-28
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Appendicectomy, Appendicitis, Appendicolith, Computerised tomogram abdomen abnormal, Laboratory test, Ultrasound abdomen abnormal
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal obstruction (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Back pain
Allergies: None
Diagnostic Lab Data: Labs drawn, abdominal ultrasound and ct scan performed. Stones found in appendix
CDC Split Type:

Write-up: Abdominal pain that proceeded to get worse into the next day. Connected with PCP, had labs drawn and ultrasound ordered. Ended up going to ER. Determined to have

Appendicitis, needed and had appendectomy on 12/29/2020.

VAERS ID: 921053 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Indiana
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	- / OT

Administered by: Private **Purchased by:** ?
Symptoms: Cardiac monitoring abnormal, Flushing, Heart rate increased, Supraventricular tachycardia, Urticaria
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201228; Test Name: heart rate; Result Unstructured Data: Test Result:180"s earlier in the day; Test Date: 20201228; Test Name: heart rate; Result Unstructured Data: Test Result:increased to 160"s
CDC Split Type: USPFIZER INC2020518671

Write-up: Patient had SVT; flushing; hives; heart rate increased to 160"s (had been 180"s earlier in the day); This is a spontaneous report from a contactable pharmacist. A 46-year-old female patient received the first dose of BNT162B2 (lot number: EK5730), via intramuscular, on 28Dec2020 at single dose for COVID-19 immunization. Facility where the most recent COVID-19 vaccine was administered was hospital. The patient was not pregnant at the time

of vaccination. The patient was not diagnosed with COVID-19 prior to vaccination. Since the vaccination, it''s unknown if the patient was tested for COVID-19. No other vaccines were received within 4 weeks prior to the COVID vaccine. The patient''s medical history and concomitant medications were not reported. The patient had SVT, flushing, hives 20 min after receiving vaccine on 28Dec2020. Patient was taken to ED and evaluated. SVT resolved. Patient sent home on heart monitor. Later that night while in bed, heart rate increased to 160''s (had been 180''s earlier in the day) and patient was admitted to hospital. Patient is a NP. Treatment received for the adverse event included cold water to face, vagal massage. The outcome of the event"Patient had SVT" was recovered on 28Dec2020 and of other events was recovering.; Sender''s Comments: A causal association between BNT162B2 and the reported events supraventricular tachycardia, flushing, hives, heart rate increased cannot be excluded based on the compatible temporal relation. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 921062 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 2020-12-01
Onset: 2020-12-01
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Fatigue, Intensive care, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020520682

Write-up: a high fever; extreme fatigue; have allergies; This is spontaneous report from a non-contactable consumer. This consumer reported similar events for eight patients. This is the first of eight reports. Only this report is serious. A patient of unspecified age and gender received bnt162b2 (BNT162B2 also reported as Pfizer version of the vaccine, lot/batch number and expiry date were not reported), via an unspecified route of administration on unknown date in Dec2020 at single dose, for Covid-19 immunisaton. The patient medical history and concomitant medications were not reported. The patient got the vaccine at a facility then had a high fever and extreme fatigue after getting the vaccine on Dec2020. The patient was admitted to the ICU. The patient had allergies, but it was unknown what the allergies are to. The outcome of events was unknown. No follow-up attempts are possible. Information on batch/Lot number can not be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2020520700 same reporter/drug/AE, different patients;US-PFIZER INC-2020520703 same reporter/drug/AE, different patients;US-PFIZER INC-2020520699 same reporter/drug/AE, different patients;US-PFIZER INC-2020520704 same reporter/drug/AE, different patients;US-PFIZER INC-2020520705 same reporter/drug/AE, different patients;US-PFIZER INC-2020520701 same reporter/drug/AE, different patients;US-PFIZER INC-2020520702 same reporter/drug/AE, different patients;US-PFIZER INC-2020520700 same reporter, drug, events, and different patients;US-PFIZER INC-2020520701 same reporter, drug, events, and different patients

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VAERS ID: 921087 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1686 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Cough, Speech disorder, Stridor
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Dementia (broad), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No

ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: lisinopril/HCTZ, albuterol MDI
Current Illness: no
Preexisting Conditions: asthma
Allergies: ibuprofen, IV contrast allergy
Diagnostic Lab Data:
CDC Split Type:

Write-up: Pt received COVID Vaccine at 1055, 1120 pt began coughing severely and could not stop, unable to speak. 1120 25mg Benadryl liquid given, Pepcid 20 mg given PO, cough worsening. 1122 second dose of Benadryl given, called for MD. Brought pt to Private room via wheelchair. Upon arrival, audible stridor noted. Epinephrine 0.3 mg IM given at 1126. IV started, placed patient on monitor and O2 via 1L NC. MD at bedside along with RT and pharmacy. 1134 Solumedrol 125 mg IV given, 1 puff of Ventolin given. Lungs clear. 1140 Coughing stopped, pt able to speak now. Vital signs: 1130 SPO2 99% Pulse 142 1135 99% pulse 106 BP 168/102 1140 sats 100% HR 93 BP 157/105 1145 sats 100% HR 97 BP 159/93 1200 sats 99% HR 103 155/97 114

VAERS ID: 921090 (history)
Form: Version 2.0
Age: 77.0
Sex: Female
Location: Minnesota
Vaccinated: 2020-12-23
Onset: 2020-12-31
Days after vaccination: 8
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Alanine aminotransferase increased, Aspartate aminotransferase increased, Brain natriuretic peptide increased, COVID-19, Dyspnoea, Fatigue, Fibrin D dimer increased, International normalised ratio decreased, SARS-CoV-2 test positive, Troponin increased
SMQs:, Cardiac failure (broad), Liver related investigations, signs and symptoms (narrow), Anaphylactic reaction (broad), Haemorrhage laboratory terms (broad), Myocardial infarction (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Apixaban, ECASA, Atorvastatin, Labetalol, Multivitamin w/minerals, Pantoprazole, Diltiazem, Meclizine
Current Illness: Mitral valve surgery (surgery planned within days)
Preexisting Conditions: HTN, GERD, A Fib, Sleep apnea on CPAP, Lactose intolerance, allergies
Allergies: Sulfa/Trimethoprim
Diagnostic Lab Data: on 1/4/2021: COVID-19 Pos ; D Dimer 243; BNP 400; Troponin 10; AST 47; ALT 42; INR 2
CDC Split Type:

Write-up: Pt vaccinated on 12/23. PCP notified that SOB and fatigue getting worse on 1/4. Unable to keep pre-op Dental work planned prior to mitral valve surgery on 1/14/2021. PCP referred her to our ED where she was diagnosed with COVID-19 and transferred to facility, which is where her surgery was planned.

VAERS ID: 921091 (history)
Form: Version 2.0
Age: 54.0
Sex: Male
Location: New Hampshire
Vaccinated: 2020-12-31
Onset: 2020-12-31
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / UNK	RA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Defaecation urgency, Hyperhidrosis, Loss of consciousness, Nausea, Road traffic accident, Syncope
SMQs:, Torsade de pointes/QT prolongation (broad), Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Accidents and injuries (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Noninfectious diarrhoea (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none related
Current Illness:
Preexisting Conditions: diabetes
Allergies: neomycin
Diagnostic Lab Data:
CDC Split Type:

Write-up: Pt received vaccination and left after 15 min. observation symptom free. He drove a short distance away from the clinical site when he felt profuse sweating and had syncope (seconds) crashing into curb. He was aroused from impact and was able to stop car. He then developed profuse nausea and sudden urge to defecate. He went to restroom. Given these events he returned to the clinical site in another vehicle. Upon arrival he denied chest pain, shortness of breath or ongoing nausea or abdominal pain. He reported his AM blood sugar was 72 and does not take insulin. No history of coronary disease or syncope. EMS was activated and assumed care.

VAERS ID: 921151 (history)
Form: Version 2.0
Age: 43.0
Sex: Female
Location: New York
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood HIV RNA below assay limit, CD4 lymphocyte percentage decreased, Condition aggravated, Endotracheal intubation, Intensive care, Mechanical ventilation, Seizure
SMQs:, Angioedema (broad), Systemic lupus erythematosus (broad), Convulsions (narrow), Acute central respiratory depression (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Respiratory failure (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: HIV DM HTN Seizure d/o Schizoaffective d/o Lipid d/o
Preexisting Conditions: As above
Allergies: Penicillin, Bactrim, Dapsone
Diagnostic Lab Data: None performed at the nursing home at the time of transfer. Last Keppra level = 19 (therapeutic range -10-40 ug/ml) , done on 12/11/20 Last CD4 count = 1123/43.2 % , HIV viral load <20; done on 6/20/20
CDC Split Type:

Write-up: The resident who was known to have seizures, and under control for many yeas with Keppra 1000 mg twice a day, on the second day after vaccination developed recurrent seizures requiring hospitalization to an intensive care unit, with intubation and mechanical ventilation until 1/5/21 (to be extubated today). She is still at the hospital.

VAERS ID: 921171 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: Virginia
Vaccinated: 2020-12-22
Onset: 2020-12-24
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Asthma, COVID-19 pneumonia, Computerised tomogram thorax abnormal, Condition aggravated, Diarrhoea, Dyspnoea, Gastrointestinal disorder, Lung consolidation, Lung opacity, PO2 decreased, SARS-CoV-2 test positive, Wheezing
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Asthma/bronchospasm (narrow), Interstitial lung disease (narrow), Pseudomembranous colitis (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypersensitivity (broad), Noninfectious diarrhoea (narrow), Respiratory failure (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Famotidine, albuterol inhaler, xopenex
Current Illness: Asthma
Preexisting Conditions: asthma
Allergies: aspirin
Diagnostic Lab Data: COVID POSITIVE;" PO2 66.2; O2 SAT 88% ROOM AIR; Severe and diffuse ground-glass, patchy, and crazy paving airspace consolidations are seen throughout the lungs; Severe, diffuse, and bilateral COVID-19 pneumonia shown on CT of chest.
CDC Split Type:

Write-up: RECEIVED VACCINE ON 12/22; ON 12/24, STARTED FEELING WEAK AND HAVING GI ISSUES WITH DIARRHEA. ON 12/27, STARTED HAVING SHORTNESS OF BREATH AND WHEEZING MORETHAN HER NORMAL WITH HER ASTHMA ILLNESS AND CAME TO ER. WAS TESTED AND FOUND TO BE COVID POSITIVE.

VAERS ID: 921175 (history)
Form: Version 2.0
Age: 77.0
Sex: Female
Location: New Jersey
Vaccinated: 2021-01-03
Onset: 2021-01-03
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1686 / 1	- / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Body temperature increased, Dyspnoea
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-05
Days after onset: 2
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: CHF, COPD, DM, heart failure, anemia, sleep apnea
Preexisting Conditions: CHF, COPD, DM, heart failure, anemia, sleep apnea
Allergies: NKA
Diagnostic Lab Data: Epi pen 0.3mg given, sent to ER
CDC Split Type:

Write-up: Resident received Covid Vaccine, noted after 30 mins with labored breathing BP 161/77, HR 116, R 38, T 101.4,

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VAERS ID: 921465 (history)
Form: Version 2.0
Age: 51.0
Sex: Female
Location: Rhode Island
Vaccinated: 2020-12-27
Onset: 2020-12-30
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Arthralgia, Back pain, Chest X-ray normal, Chills, Computerised tomogram normal, Computerised tomogram spine, Ear pain, Headache, Laboratory test, Myalgia, Nausea, Pain, Pain in extremity, Photophobia, Purpura, Pyrexia, Rash, Vomiting
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Acute pancreatitis (broad), Haemorrhage terms (excl laboratory terms) (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Noninfectious meningitis (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Glaucoma (broad), Corneal disorders (broad), Eosinophilic pneumonia (broad), Retinal disorders (broad), Hypersensitivity (narrow), Arthritis (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: 51-year-old female with history of intermittent asthma presented to the ED with 1 week of intermittent fevers, myalgias, arthralgias and headache. Patient reports receiving first dose of moderna vaccine last week. She initially developed arm soreness followed by chills and body aches. Subsequently developed frontal headache, photophobia, back pain, nausea and vomiting. She was seen in the ER on 1/1/21, when her work-up including labs, CT spine, chest x-ray were negative therefore she was discharged home. She continued to have symptoms and also developed bilateral intermittent ear pain. She also developed rash in her extremities and torso. Rash is pruritic but not painful. reports ongoing history of neck pain for which she sees PT. Denies sore throat, cough, chest pain or shortness of breath.

VAERS ID: 921557 (history)
Form: Version 2.0
Age: 74.0
Sex: Male
Location: Minnesota
Vaccinated: 2021-01-03
Onset: 2021-01-04
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Asthenia, Blood pressure decreased, Body temperature increased, Chest X-ray normal, Heart rate increased, Hyponatraemia, Metabolic function test abnormal, Urine analysis normal
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Hyponatraemia/SIADH (narrow), Chronic kidney disease (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: cholecalciferol, iron, oxycodone, gabapentin

Current Illness:
Preexisting Conditions: obesity, GERD, allergic rhinitis, neuropathy, insomia, ataxia, hyperlipidemia
Allergies: cosopt, weeds, soaps
Diagnostic Lab Data: Chest X-Ray--unremarkable, UA--unremarkable, BMP--mild hyponatremia
CDC Split Type:

Write-up: Began experiencing increased temp of 101.4 on 1/4/2021 at 0701. Temp did not resolve with the use of Tylenol. HR increased to $g100 and BP was decreasing below baseline. Increased weakness also noted. Temp increased to 102.9 on 1/4/2021 at 2220. Transferred from SNF to ER for evaluation.

VAERS ID: 921641 (history)
Form: Version 2.0
Age: 17.0
Sex: Female
Location: New York
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Acute respiratory failure, COVID-19, Hypoxia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Eosinophilic pneumonia (broad), Hypersensitivity (broad), Respiratory failure (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: acetaminophen 160 mg/5 mL (5 mL) oral solution albuterol sulfate 2.5

mg/3 mL (0.083 %) solution for nebulization budesonide 0.5 mg/2 mL suspension for nebulization Certavite-Antioxidant 18 mg-400 mcg tablet ClearLax 17 gram/dose oral powder
Current Illness: Right lung atelectasis/infiltrate with right mediastinal shift on chest x-ray (12/25) that was treated with steroids and antibiotics
Preexisting Conditions: Chronic respiratory Failure Dependence on mechanical ventilation Seizure disorder Spastic quadriplegic cerebral palsy Encephalopathy
Allergies: NKA
Diagnostic Lab Data: 1/5/21 COVID PCR result: Positive
CDC Split Type:

Write-up: Administered first dose of COVID19 vaccine at 1:29pm on 1/4/21. At approximately 11:00pm resident exhibited acute respiratory decompensation with very limited air entry and hypoxemia. Patient received Benadryl, steroids, epinephrine, and Duoneb without improvement. Resident was referred to the emergency room and found to be COVID positive. No fever or rash were reported.

VAERS ID: 921786 (history)
Form: Version 2.0
Age: 75.0
Sex: Male
Location: Michigan
Vaccinated: 2021-01-03
Onset: 2021-01-03
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Tremor
SMQs:, Neuroleptic malignant syndrome (broad), Parkinson-like events (broad), Noninfectious encephalopathy/delirium (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Metformin, Keppra, Eliquis, Metoprolol Succinate, Mesalamine, Colace , bupropin ER and Ferrous Sulfate given at 10am on 1-3-21
Current Illness:
Preexisting Conditions: A-fib, HTN, Type 2 DM, Dementia and hyperlipidemia
Allergies: NKA
Diagnostic Lab Data:

CDC Split Type:

Write-up: tremors resident sent hospital facility.

VAERS ID: 921796 (history)
Form: Version 2.0
Age: 49.0
Sex: Female
Location: Tennessee
Vaccinated: 2021-01-02
Onset: 2021-01-02
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / IM

Administered by: Public **Purchased by:** ?
Symptoms: Cardiovascular evaluation, Chest X-ray, Chest discomfort, Chest pain, Dehydration, Feeling abnormal, Headache, Laboratory test, Nausea, SARS-CoV-2 test negative, Tinnitus, Urine analysis, Vaccination complication, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Dementia (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hearing impairment (narrow), Dehydration (narrow), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Wellbutrin, Microbid
Current Illness: no
Preexisting Conditions: ESBL
Allergies: Buspar
Diagnostic Lab Data: heart work up, COVID (negative) labs, chest xray, urine
CDC Split Type:

Write-up: Started out vague. Started with headache at the base of her head and then it felt like a web that covered her entire head. By 10:00 she did not feel good. Went to sleep instantly. Slept for about an hour. Began to have nausea. Sunday headache got worse. Started ringing , buzzing sound in her head. Took Zofran because of nausea. Sunday night chest discomfort. Monday had horrible chest pain, headache was horrible, then vomiting. went to ER, doctor felt it was a reaction to the vaccine. was give medicine for nausea and Decadron for the reaction. Gave fluids for dehydration. Got medicine for headache. Keep

taking Zofran for nausea. Still has headache. Has appointment with pcp in the morning.

VAERS ID: 921829 (history)
Form: Version 2.0
Age: 81.0
Sex: Male
Location: Minnesota
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Chest X-ray abnormal, Diarrhoea, Hypoxia, Lung infiltration, Nausea, Pyrexia, Vomiting, White blood cell count normal
SMQs:, Acute pancreatitis (broad), Asthma/bronchospasm (broad), Interstitial lung disease (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Noninfectious diarrhoea (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Acetaminophen, Aspercreme Lidocaine Liquid 4 %, Biotene Dry Mouth Liquid (Mouthwashes, ,Calcium Antacid Tablet Chewable 500 MG (CalciumCarbonate Antacid) ,Carbidopa-Levodopa Tablet, Coumadin Tablet, Fiber formula, furosemide, gabapentin,Ins
Current Illness: COVID and pneumonia
Preexisting Conditions: CHF, COPD, parkinsons, diabetes, hyperlipidemia, obstructive sleep apnea, hypertension, dilated cardiomyopathy, hyperparathyroidism
Allergies: Lisinopril
Diagnostic Lab Data: Chest x-ray, CBC,
CDC Split Type:

Write-up: Received COVID-19 Moderna vaccine on 12/28. Developed nausea, vomiting, diarrhea, fever, and hypoxia at facility the day follow vaccine administration (12/29). He was sent to the hospital on 12/29 and admitted for post vaccine fever where he had a chest x-ray

that showed infiltrates but WBC count was normal and fever resolved upon admission to hospital. Provider documented "expected reaction to vaccination in a patient with previous COVID exposure". He returned to the facility on 12/30.

VAERS ID: 921920 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Michigan
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	UN / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, Oxygen saturation decreased, Posture abnormal, Somnolence
SMQs:, Anaphylactic reaction (broad), Anticholinergic syndrome (broad), Dementia (broad), Dystonia (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Respiratory failure (broad), Hypoglycaemia (broad), Infective pneumonia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None known
Current Illness: None listed
Preexisting Conditions: None listed
Allergies: None listed
Diagnostic Lab Data: Decrease in O2 levels
CDC Split Type:

Write-up: Within 10 minutes of receiving the vaccine patient began to look sleepy and started to gasp for breath. She called for help and slouched over in a chair and was moved to lay on the floor. Her feet were elevated and 911 was called. Epinephrine was given, patient responded by being able to gasp for breath and her face returned to a more natural state. Within 5 min she began to struggle to breath and epi was given again. Some improvement did occur. EMS arrived and she was taken to the hospital.

VAERS ID: 921989 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Dizziness, Dyspnoea, Erythema, Obstructive airways disorder, Swelling, Swelling face
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Asthma/bronchospasm (broad), Anticholinergic syndrome (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: High cholesterol Frequent UTIs
Preexisting Conditions:
Allergies: None
Diagnostic Lab Data:
CDC Split Type:

Write-up: Anaphylactic reaction (swelling and redness of face and torso, shortness of breath, constriction of airway and dizziness)

VAERS ID: 922118 (history)
Form: Version 2.0

Age: 59.0
Sex: Male
Location: Georgia
Vaccinated: 2020-12-31
Onset: 2021-01-03
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood pressure increased, Feeling jittery, Insomnia
SMQs:, Neuroleptic malignant syndrome (broad), Hypertension (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Edarbyclor, Allopurinol, Apple Cider
Current Illness: no
Preexisting Conditions: High blood pressure
Allergies: Lisinopril
Diagnostic Lab Data: no
CDC Split Type:

Write-up: After three days, couldn"t sleep the whole night. The next day, went to work, came home felt jittery. Close to midnight bp 200/90. Took Clonidine 0.1 and went to ER, bp 180/90. waited for almost two hours bp came down to 141/80. Today, bp is back to normal. Took sleeping medication Zzzquil to go to sleep.

VAERS ID: 922177 (history)
Form: Version 2.0
Age: 28.0
Sex: Male
Location: Connecticut
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J2OA / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Alanine aminotransferase increased, Borrelia test negative, CSF culture negative, CSF glucose normal, CSF mononuclear cell count increased, CSF polymorphonuclear cell count increased, CSF protein normal, CSF red blood cell count positive, CSF virus no organisms observed, CSF white blood cell count increased, Cryptococcus test, HIV test negative, Headache, Hepatitis A virus test, Hepatitis B test negative, Hepatitis C test negative, Herpes simplex test negative, Influenza virus test negative, Magnetic resonance imaging brain normal, Meningitis aseptic, Mononucleosis heterophile test negative, Polymerase chain reaction, Pyrexia, SARS-CoV-2 test negative, Staphylococcus test positive, Treponema test negative, Varicella virus test negative
SMQs:, Liver related investigations, signs and symptoms (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Noninfectious meningitis (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none
Preexisting Conditions: none
Allergies: none
Diagnostic Lab Data: Done on 12/31/ (ALT is 124), Hepatitis A and B, and C are negative. Mono negative, COVID 19 PCR is negative Influenza PCR negative; Blood culture x 1 bottle + for Staph hemolyticus and Staph hominis 1/1/2021; Lumbar tap showed elevated WBC of 23, 76% polys and 24% mononuclear, 25 RBC. Glucose and protein are normal. Culture negative. CSF PCR viral panel is negative, RPR negative, Lyme serology is negative, Crypto Antigen is negative. 1/2/2021 - HIV serology is negative 1/5/2021 - MRI of the brain is normal
CDC Split Type:

Write-up: Aseptic meningitis, prolonged fever for more than a week, headache, elevated transaminase (ALT is 124). Lumbar tap showed elevated WBC of 23, 76% polys and 24% mononuclear, 25 RBC. Glucose and protein are normal. CSF PCR viral panel is negative. Patient was initially given Acyclovir and was stopped when HSV and VZV PCR were negative. He was given vancomycin IV for Gram positive bacteremia which was later stopped because it was deemed a contaminant.

VAERS ID: 922218 (history)
Form: Version 2.0
Age: 41.0
Sex: Female

Location: Colorado
Vaccinated: 2020-12-23
Onset: 2020-12-26
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Chills, Cold sweat, Dizziness, Feeling cold, Headache, Hypotension, Pain, Peripheral coldness, Pyrexia, Septic shock, White blood cell count increased
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Toxic-septic shock conditions (narrow), Vestibular disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (broad), Sepsis (narrow), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Entyvio
Current Illness:
Preexisting Conditions: Crohns disease
Allergies: N/A
Diagnostic Lab Data:
CDC Split Type:

Write-up: 3 Days after the covid vaccine. I Started to have these symptoms around 1am: fever, chills, body aches, cold sweats. I took ibuprofen fell asleep. Around 330am. Woke up with dizziness, headache, chills. Took tylenol. Fell asleep body was extremely cold esp hands and feet. Woke up around 7am. Still had severe body aches and was extremely dizzy, headache worsening. Took ibuprofen. Woke up around noon. Extremely dizzy with chills and body aches headache still severe. Took tylenol. I was not getting better. went to hospital . It was found I had extremely high wbc and had extremely low blood pressure. Diagnosed as septic shock.

VAERS ID: 923540 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Unknown

Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / UNK	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Dizziness, Dysphagia, Dyspnoea, Erythema, Headache, Lip swelling, Peripheral swelling, Pharyngeal swelling, Stridor, Tachypnoea
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Angioedema (narrow), Asthma/bronchospasm (broad), Anticholinergic syndrome (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Anaphylaxis Narrative: 12/22 received COVID-19 vaccine at 1209 and developed SOB at 12:15. Took her own albuterol inhaler without relief. Transported to ED. PE: red hands with swelling, throat and lip swelling with difficulty swallowing. Later developed headache and dizziness then tachypnea and stridor. Meds given - See section 5 PLUS epinephrine IM and infusion @ 0.05 mcg/kg/min, Alb and ipratropium negs, racemic epi nebs. Admitted to the hospital on 12/22 and still hospitalized at the time of this report on 12/23. She remains on an epinephrine drip and was given methylprednisolone 125 mg IV x 2. No previous history of anaphylaxis. History of Reye"s syndrome as a child when given aspirin.

VAERS ID: 923554 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 2020-12-23
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LL / IM

Administered by: Other **Purchased by:** ?
Symptoms: Agitation, Anaphylactic reaction, Hypotension, Rash, Sedation
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hostility/aggression (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Agitation, Sedation, Anaphylaxis, Rash & HYPOtension

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VAERS ID: 919843 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-27

Onset: 2020-12-28
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Epilepsy
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Generalised convulsive seizures following immunisation (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Dementia
Allergies:
Diagnostic Lab Data:
CDC Split Type: DEPFIZER INC2020517434

Write-up: generalised epileptic fit; This is a spontaneous report from a contactable physician. A 92-year-old female patient started to received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 27Dec2020 as single dose for covid-19 immunization. Medical history included dementia from an unknown date. The patient''s concomitant medications were not reported. The patient experienced generalised epileptic fit on 28Dec2020, which was serious per noted hospitalization. The patient was hospitalized for generalised epileptic fit from 28Dec2020 to 29Dec2020. The outcome of generalised epileptic fit was recovered on 29Dec2020; the patient recovered to her previous level. No follow-up attempts are possible; information about batch/lot number cannot be obtained.; Sender''s Comments: The reported information is limited and does not allow for a meaningful assessment of the case .The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 919853 (history)
Form: Version 2.0
Age:

Sex: Female
Location: Foreign
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic shock, Angioedema, Guillain-Barre syndrome, Muscle spasms, Myelitis transverse, Paraesthesia, Paralysis, Rash, Respiratory distress, Seizure
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Peripheral neuropathy (narrow), Systemic lupus erythematosus (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Convulsions (narrow), Dystonia (broad), Oropharyngeal allergic conditions (narrow), Acute central respiratory depression (broad), Guillain-Barre syndrome (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Demyelination (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Generalised convulsive seizures following immunisation (narrow), Hypersensitivity (narrow), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: MXPFIZER INC2020521244

Write-up: rash; spasms; paresthesia/lower limb paresthesia; Arms and legs paralysis; Guillain Barre syndrome; angioedema; impaired respiratory distress; anaphylactic shock; transverse myelitis; post-vaccine application seizures; This is a spontaneous report from a contactable physician and consumer. A 3-decade-old female patient received first dose of bnt162b2 (BNT162B2, lot number was not reported), intramuscular on 30Dec2020 09:30 at a single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient had a reaction to the vaccine, she was in serious condition. On 30Dec2020 09:50, 20 minutes after the application, the patient experienced rash. After that, the patient developed angioedema with impaired respiratory distress, spasms and

paresthesia. The patient was hospitalized and required transfer to the Intensive care unit of the hospital. The reporter also stated that it was probably compatible with an adverse event for the vaccination like Guillain-Barre syndrome. The patient also developed anaphylactic shock, lower limb paresthesia and post-vaccine application seizures, arms and legs paralysis, frequent convulsions, that evolved to transverse myelitis, valued by the doctor that suggested intensive therapy of third level. The events caused hospitalization and the outcome of the events was unknown.; Sender''s Comments: The reporter provided only limited and anecdotal information and the rationale for suspecting a causal relationship between the events and BNT162b2 is not evident from the data provided apart from an assumed chronological association. Due to the lack of critical information (e.g. complete patient''s demographics, medical history, concomitant medications, diagnostic work-up for the reported clinical conditions) an independent clinical evaluation is impossible at this time.

VAERS ID: 920906 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-16
Onset: 2020-12-17
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Epilepsy, Generalised tonic-clonic seizure
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: CALCEOS; ; ; ; ; ; ; CASSIA SENNA; ; ; TEGRETOL; ; ; ; EPILIM CHRONO; CO-AMOXICLAV [AMOXICILLIN;CLAVULANATE POTASSIUM]
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Epilepsy
Allergies:

Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2020517522

Write-up: Tonic-clonic seizures x 3; Tonic-clonic seizures x 3; This is a spontaneous report from a contactable pharmacist downloaded from the Medicines Agency (MA) Regulatory Authority-WEB [GB-MHRA-EYC 00235725]. Safety report unique identifier [GB-MHRA-ADR 24544683]. A 62-year-old female patient received BNT162B2 (lot number unknown), intramuscular on 16Dec2020 at single dose for COVID-19 immunization. Medical history included epilepsy from an unknown date and unknown if ongoing. Concomitant medication included calcium carbonate, colecalciferol (CALCEOS), lansoprazole, amoxicillin, cyclizine, alendronic acid, ferrous fumarate, clopidogrel, cassia senna, atorvastatin, paracetamol, carbamazepine (TEGRETOL), trimethoprim, diazepam, atenolol, valproate sodium, valproic acid (EPILIM CHRONO), amoxicillin, clavulanate potassium (CO-AMOXICLAV), all of which were taken from unspecified dates for unspecified indications. It was reported that a nursing home patient with past medical history of epilepsy. He had three tonic clonic seizures (tonic-clonic seizures x 3) the day after receiving BNT162B2 vaccine (17Dec2020), resolved by administration of rectal diazepam. The patient was hospitalized for the events. The patient had no known drug allergies. The outcome of the events was resolved on 17Dec2020. No follow-up attempts are possible. No further information expected, information on batch number cannot be obtained.; Sender''s Comments: This 62-year-old female patient had medical history included ongoing epilepsy. The reported event of three tonic clonic seizures was more likely due to underlying condition, and less likely causally related to the use of BNT162B2 on 16Dec2020 at single dose for COVID-19 immunization. The case will be reassessed should additional information become available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 920907 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 0000-00-00
Onset: 2020-12-16
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Cerebral infarction
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; CO-CODAMOL; TRIMBOW; ; ; ; EVACAL D3; ; ;
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2020517463

Write-up: Partial Anterior Circulation Infarct; This is a spontaneous report from a contactable healthcare professional downloaded from the Medicines Agency (MA) WEB (GB-MHRA-EYC 00235731 and GB-MHRA-ADR 24544793). A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, on an unspecified date at a single dose for COVID-19 vaccination. The patient"s medical history was not reported. Concomitant medications included levothyroxine (MANUFACTURER UNKNOWN), lidocaine (MANUFACTURER UNKNOWN), ferrous fumarate (MANUFACTURER UNKNOWN), codeine phosphate, paracetamol (CO-CODAMOL), beclometasone dipropionate, formoterol fumarate, glycopyrronium bromide (TRIMBOW), salbutamol (MANUFACTURER UNKNOWN), amitriptyline (MANUFACTURER UNKNOWN), simvastatin (MANUFACTURER UNKNOWN), calcium carbonate, colecalciferol (EVACAL D3), metformin (MANUFACTURER UNKNOWN), ramipril (MANUFACTURER UNKNOWN), and gabapentin (MANUFACTURER UNKNOWN). The patient experienced partial anterior circulation infarct on 16Dec2020, which caused hospitalization. The clinical outcome of partial anterior circulation infarct was recovered with sequel on an unspecified date. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 920911 (history)
Form: Version 2.0
Age: 83.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	E50553 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Transient ischaemic attack
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Embolic and

thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ADCAL [CALCIUM CARBONATE]; ; ; ; ; PERINDOPRIL TERT-BUTYLAMINE
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Arthroplasty of hip (Cement Used); Degenerative joint disease (Acromioclavicular joint); Femoral neck fracture (Closed fracture of neck of femur); Osteoporosis
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2020517620

Write-up: Transient ischaemic attack; This is a spontaneous report from contactable physician downloaded from the Medicines Agency (MA) WEB regulatory authority or other manufacturer number GB-MHRA-TPP21563188C6255921YC1608215659743, Safety Report Unique Identifier GB-MHRA-ADR 24544363 received via regulatory authority. An 83-year-old female patient receive BNT162B2 (COVID-19 MRNA VACCINE BIONTECH, batch/lot number E50553, with unknown expiration date), Multidose vial, parenteral on 17Dec2020 at 0.3 mL single for covid-19 immunisation. Medical history included degenerative joint disease (Acromioclavicular joint) from an unknown date and unknown if ongoing; hip arthroplasty (Cement Used) from an unknown date and unknown if ongoing, femoral neck fracture (Closed fracture of neck of femur) from an unknown date and unknown if ongoing, osteoporosis from an unknown date and unknown if ongoing. Concomitant medication included alendronic acid from 13Mar2020, calcium carbonate (ADCAL) from 27May2020, citalopram (CITALOPRAM) from 15Dec2015, atorvastatin from 15Dec2015, propranolol from 21Nov2017, apixaban from 06Aug2019, perindopril erbumine from 15Dec2015. The patient experienced transient ischaemic attack (hospitalization, medically significant) on 17Dec2020 with outcome of recovered on 17Dec2020. Admitted with transient ischaemic attack symptoms hours after her Pfizer vaccine. No follow-up attempts possible. No further information expected.; Sender"s Comments: This 83-year-old female patient had ongoing medical history and was on multiple concomitant medications. Based on information available, the reported event transient ischaemic attack was most likely an intercurrent disease, and unlikely causally related to the immunization with BNT162B2 (COVID-19 MRNA VACCINE BIONTECH). The case will be reassessed should additional information become available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 920917 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-16
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood test, Dysphagia, Hypersensitivity, Lip swelling, Malaise, SARS-CoV-2 test, Swelling face
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: INFLUENZA VIRUS; ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Adverse reaction; Comments: Patient has not had symptoms associated with COVID-19 Patient is not enrolled in clinical trial
Allergies:
Diagnostic Lab Data: Test Name: blood test for allergic response; Result Unstructured Data: Test Result:unknown result; Test Date: 20201024; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:Yes - Positive COVID-19 test
CDC Split Type: GBPFIZER INC2020517804

Write-up: Swelling lips; Swallowing difficult; Feeling unwell; Allergy; Swelling face; This is a spontaneous report from a contactable consumer downloaded from the Medicines Agency (MA) Regulatory Authority-WEB, Regulatory Authority number: GB-MHRA-ADR 24544024 and GB-MHRA-WEBCOVID-20201216194551. An 87-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) (batch/lot number: unknown), via an unspecified route of administration on 15Dec2020 at single dose for COVID-19 immunisation. Medical history included bad reaction to mosquitos from an unknown date and unknown if

ongoing. Concomitant medication included influenza vaccine (INFLUENZA VIRUS), fentanyl, paracetamol, ramipril, pregabalin. The patient experienced swelling lips (hospitalization, medically significant) on 16Dec2020, swallowing difficult (hospitalization, medically significant) on 16Dec2020, feeling unwell (hospitalization, medically significant) on 16Dec2020, allergy (hospitalization, medically significant) on 16Dec2020, swelling face (hospitalization, medically significant) on 16Dec2020. The patient underwent lab tests and procedures which included the result of blood test for allergic response was unknown result in unknown date; the result of COVID-19 virus test was yes - positive covid-19 test on 24Oct2020. Patient had not had symptoms associated with COVID-19. Patient was not enrolled in clinical trial. Patient Woke up morning after vaccine with a swollen face which worsened with time spreading to lips and Felt very unwell and had difficulty swallowing. Ambulance called and patient admitted to hospital for hydrocortisone and antihistamine drip. Breathing not affected. Patient had not tested positive for COVID-19 since having the vaccine. The outcome of the event allergy was not recovered. The outcome of other events was unknown. No follow-up attempts possible. No further information expected. Batch/lot number cannot be obtained.

VAERS ID: 920921 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-13
Onset: 2020-12-13
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood test, Body temperature, Chills, Dyspnoea, Feeling hot, Feeling jittery, Flushing, Heart rate, Medical observation, Pyrexia, Rash, Rash macular, Tachycardia
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions: Comments: No medical history Patient has not had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial Patient is not pregnant
Allergies:
Diagnostic Lab Data: Test Date: 20201213; Test Name: Blood test; Result Unstructured Data: Test Result:slightly raised c-reactive protein; Test Date: 20201213; Test Name: Blood test; Result Unstructured Data: Test Result:slightly raised white cell count; Test Date: 20201213; Test Name: Body temperature; Result Unstructured Data: Test Result:37.8 at 5:53pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:96 on admission; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:increasing to 140 on exertion - around 5pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:still raised at 7:30pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:105 - at rest, around 5pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:138 at 5pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:raised at 100 - around 5pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:138 on minor exertion - 7:30pm; Test Date: 20201213; Test Name: Pulse rate; Result Unstructured Data: Test Result:over 90 at 4pm; Test Date: 20201213; Test Name: observation; Result Unstructured Data: Test Result:observations okay until 4pm
CDC Split Type: GBPFIZER INC2020517621

Write-up: Body temperature 37.8; Feeling hot; Shivering; Shortness of breath; Tachycardia; Macular rash; Jitteriness; Rash; Flushing; This is a spontaneous report from contactable physician downloaded from the Medicines Agency (MA) Regulatory Authority-WEB with regulatory authority number GB-MHRA-WEBCOVID-20201217072238, Safety Report Unique Identifier GB-MHRA-ADR 24544078. A 34-year-old female patient (not pregnant) received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, batch/lot number unknown, with unknown expiration date), via an unspecified route of administration on 13Dec2020 14:45 at single dose for COVID-19 vaccination. The patient"s medical history and concomitant medications were not reported. Patient has not had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Patient was not enrolled in clinical trial. The patient was a healthy female with no allergies. The patient experienced feeling hot on 13Dec2020, shivering on 13Dec2020, shortness of breath on 13Dec2020, tachycardia on 13Dec2020, macular rash on 13Dec2020, Jitteriness on 13Dec2020, rash on 13Dec2020, flushing on 13Dec2020. All led to hospitalization. It was reported that after 14:45 vaccinated, the patient felt hot, flushed, jittery, shivery about 5 minutes later. Developed a macular rash within 5-10 minutes, looked flushed. Observations OK until 16:00 on 13Dec2020, developed mild tachycardia. Pulse over 90 at 16:00 on 13Dec2020 at this point, rash lightened and looked less flushed at this point. 17:00 on 13Dec2020, pulse 138, increasing on minor exertion. Slightly shortness of breath at this time. No swelling at any time. Pulse continued to be raised 100, 105 at rest, increasing to 140 on any exertion around 17:00 on 13Dec2020. Ambulance called at 17:15. Body temperature 37.8 at 17:53 on 13Dec2020. Felt a bit better by 19:30, but pulse still raised on 13Dec2020 and going higher to 138 on minor exertion at 19:30. Ambulance arrived at 20:20. On admission, pulse was 96 on 13Dec2020. Kept in emergency department (ED) for 3 hours. Patient has not tested positive for coronavirus disease (COVID-19) since having the vaccine. Bloods showed slightly raised c-reactive protein and white cell count on 13Dec2020. The outcome of the event tachycardia and rash was recovered on 13Dec2020. The outcome of the other events was unknown. No follow-up attempts are possible. Information about lot/batch number cannot be obtained.

VAERS ID: 920941 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-15
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	NOT KNOWN / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Headache, Heart rate, Hot flush, Paraesthesia oral, SARS-CoV-2 test, Tachycardia, Urticaria
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Chest infection (2 more incidences in Jul2020 and Aug2020); Chest pain; Chronic fatigue (Long Covid and chronic fatigue since positive diagnosis of covid in March.); COVID-19; Pneumonia; Secondary infection (pneumonia); Urinary tract infection
Allergies:
Diagnostic Lab Data: Test Date: 20201215; Test Name: Heart rate; Result Unstructured Data: Test Result:increased; Test Date: 20200312; Test Name: COVID-19 virus test; Test Result: Positive ; Comments: Yes - Positive COVID-19 test
CDC Split Type: GBPFIZER INC2020517592

Write-up: Tachycardia; Tingling tongue; Headache; Hives; Hot flushes; This is a spontaneous report from a contactable other healthcare professional downloaded from the Agency Regulatory Authority-WEB [GB-MHRA-WEBCOVID-20201218141243]. other case identifier

number: GB-MHRA-ADR 24544843. A 23-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 15Dec2020 at single dose for COVID-19 immunization. Medical history included secondary infection from Apr2020 to an unknown date (pneumonia), Chest infection from Jun2020 to Jun2020 (2 more incidences in Jul2020 and Aug2020), Chronic fatigue from Mar2020 to an unknown date (Long Covid and chronic fatigue since positive diagnosis of covid in March, on 12Mar2020), chest pain and urinary tract infection. Concomitant medication included influenza vaccine (INFLUENZA VACCINE) for immunization, amitriptyline for chest pain, nitrofurantoin for urinary tract infection, desogestrel for oral contraception. The patient experienced tachycardia, tingling tongue, headache, hives, hot flushes, all on 15Dec2020. Tingling tongue, 3 Minutes after vaccine was delivered the patient became increasingly hot and her chest broke out in hives which then travelled onto her neck and left arm. Subsequently a frontal headache formed. Increased heart rate and tachycardia was noticed in accident and emergency. The patient took loratadine 10mg about 20 minutes after reaction. The patient was given 40mg steroids prednisolone and 4mg chlorphenamine in accident and emergency and was given supply of 3 days steroids 40mg and antihistamine to take 4-6 hourly until no longer required. Serious criteria reported for all events as hospitalization. The patient underwent lab tests and procedures which included heart rate: increased on 15Dec2020, COVID-19 virus test: positive on 12Mar2020. The patient had not had symptoms associated with COVID-19. Patient was not enrolled in clinical trial. Patient was not pregnant. The outcome of events tachycardia was recovered on 18Dec2020. The outcome of event tingling tongue and frontal headache was recovered on 16Dec2020. The outcome of hives was recovering. The outcome of event hot flushes was not recovered. No follow-up attempts are possible, batch/lot number cannot be obtained. No further information is expected.

VAERS ID: 922299 (history)
Form: Version 2.0
Age: 58.0
Sex: Female
Location: Massachusetts
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Blood pressure increased, Heart rate increased, Hypoaesthesia, Hypoaesthesia oral, Laboratory test, Sensation of foreign body, Tremor
SMQs:, Anaphylactic reaction (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Hypertension (narrow), Hypoglycaemia (broad), Dehydration (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: prilosec 20mg q hs pseudofed 1 tablet at 7:15 am and 4:15pm that day. lo loestrin fe 1 tablet a day (days 1-26) off 2 days
Current Illness: cold symptoms that started 12/15/20. covid tested 12/16/20 negative. cold symptoms were diminished on 12/23/20 but still had slight runny nose. (used pseudofed 30mg tab)
Preexisting Conditions: mild asthma
Allergies: anaphylaxtis to cipro- see ER visit 4/2012. throat swelling, facial and eye swelling , head to toe hives. lasted 5 days with prednisone taper, benedryl and pepcid.
Diagnostic Lab Data: md drew labs in ER
CDC Split Type:

Write-up: I was instructed to stay for 30min as i have been anaphylatic to cipro in past. at 30min was told i could leave. while driving home on rt 91 my cheekbones became numb. then slowly a few min later my cheeks became numb. a few min later my lips became numb. as i was driving off exit to rt 5 in longmeadow i developed a lump in my throat. i turned around at top of exit and went back to highway to go to ER. this was approx 1645-1650. i went to ER arrived approx 1655. i was shaking. my bp and pulse were elevated. no tingling or swelling in my face. nurse checked my pupils and my smile and were wnl. no history of bells palsy. i received iv fluids, solucortef 125mg ivp, pepcid 20mg ivp, and benedryl 25mg ivp approx 1840pm. approx 45 min after solucortef numbness better but not gone. it started to come back a little more before discharge, which i let md know. she discharged me with scripts for epi-pen, prednisone, and OTC pepcid and benedryl. follow up with my pcp"s office in am 12/24 at 10am with his NP. total time with facial numbness/lip numbness 29 hours.

VAERS ID: 922673 (history)
Form: Version 2.0
Age: 67.0
Sex: Female
Location: Minnesota
Vaccinated: 2020-12-19
Onset: 2021-01-02
Days after vaccination: 14
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	ELO140 / UNK	LA / SYR

Administered by: Work **Purchased by:** ?
Symptoms: Angiogram, Blood test, Computerised tomogram, Computerised tomogram abdomen, Echocardiogram, Electrocardiogram, Lacunar stroke, Magnetic resonance imaging, Paraesthesia, Urine analysis

SMQs:, Peripheral neuropathy (broad), Ischaemic central nervous system vascular conditions (narrow), Guillain-Barre syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: No prescription medications Dietary supplements: Vitamin D3, Magnesium, Essential Fatty Acids, Glucosamine and Chondroitin
Current Illness: None
Preexisting Conditions: None
Allergies: Doxycycline allergy
Diagnostic Lab Data: January 2: CT scan, EKG, MRI (3 different studies), CT/angio, and lots of blood work January 3: Echocardiogram January 5: CT scan of abdomen, urinalysis and lots more blood work I will be wearing a Zia patch for two weeks after it arrives in the mail.
CDC Split Type:

Write-up: I woke up with tingling in my right hand and arm, my right side, and down my right leg. I went to the ER at Hospital, and was confirmed to have had a stroke in my left thalmus.

VAERS ID: 922767 (history)
Form: Version 2.0
Age: 87.0
Sex: Female
Location: Tennessee
Vaccinated: 2021-01-02
Onset: 2021-01-03
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	AR / IM

Administered by: Public **Purchased by:** ?
Symptoms: Anal incontinence, Asthenia, Confusional state
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Noninfectious diarrhoea (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: unknown
Diagnostic Lab Data: Patient was transported to Medical Center on 1/3/2021 and is currently still in the hospital as of today. Patients so reports they have ruled out heart attack or stroke. Her son reports they will be doing a stress test today.
CDC Split Type:

Write-up: Patient woke on 1/3/2021 weak having uncontroled bowels and off and on confusion.

VAERS ID: 922990 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: Nevada
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / UNK	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Electrocardiogram, Immediate post-injection reaction
SMQs:, Anticholinergic syndrome (broad), Vestibular disorders (broad), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: NKDA
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data: EKG (12/29/20)
CDC Split Type:

Write-up: The patient received the vaccine indicated above. Immediately following vaccination the patient states that they began feeling lightheaded and dizzy.

VAERS ID: 923000 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-17
Onset: 2020-12-26
Days after vaccination: 9
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain lower, Appendicectomy, Appendicitis, Computerised tomogram abnormal, Decreased appetite, Laparoscopic surgery, White blood cell count increased
SMQs:, Neuroleptic malignant syndrome (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Valtrex Wellbutrin Abilify Fish Oil Multi-Vitamin Biotin
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: CT Scan showed acute appendicitis resulting in a laparoscopic appendectomy. Elevated WBCs.
CDC Split Type:

Write-up: Severe right lower quadrant pain, anorexia over 12 hours. Went to the emergency department. Lab results showed elevated WBC and CT scan showed acute appendicitis. Admitted for urgent surgery: laparoscopic appendectomy. Was hospitalized from 12/26/20-

12/28/20.

VAERS ID: 923459 (history)
Form: Version 2.0
Age: 49.0
Sex: Male
Location: Oklahoma
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Flushing, Hypotension, Pyrexia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: allopurinol 100mg QD, Dilantin 30mg 2 cap QD, escitalopram 10mg QD, Zetia 10mg QD, loratadine 10mg QD, meloxicam 7.5mg QD, omeprazole 20mg QD, phenytoin EX 100mg 3 tabs QHS, phenytoin EX 100mg 2 tabs QAM, polymyxin b/trimethoprim solution 1
Current Illness: None
Preexisting Conditions: gout, optic nerve problem, dry eye syndrome, seizure disorder, depression, blepharitis, intellectual disability, Down syndrome, gingivitis, psychosis, myopia, seborrheic dermatitis, GERD, hypokalemia
Allergies: No known allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: patient was noted to be flushed by residential home staff on 12/30/2020, approximately 2.5hrs after vaccination, fever present at that time. Prior to COVID19 vaccine administration, this patient did have exposure and was close contact with a known case of COVID19 in a residential care employee. Patient was taken to hospital for evaluation of febrile status, had positive COVID19 test at that time, and reported hypotension per residential care

staff.

VAERS ID: 923510 (history)
Form: Version 2.0
Age: 85.0
Sex: Male
Location: Tennessee
Vaccinated: 2021-01-04
Onset: 2021-01-05
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Anaphylactic shock, Dyspnoea, Erythema, Injection site erythema, Injection site swelling, Peripheral swelling, Rash
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Angioedema (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: UNKNOWN
Current Illness: UNKNOWN
Preexisting Conditions: UNKNOWN
Allergies: PCN
Diagnostic Lab Data: UNKNOWN
CDC Split Type:

Write-up: PATIENT SPOUSE REPORTS THAT PATIENT RECEIVED VACCINE ON 1/4/2021 AND ON 1/5/2021 PATIENT''S ARM BEGAN TO TURN RED AND SWELL AT THE INJECTION SITE. THE SWELLING AND REDNESS BEGAN TO GO DOWN HIS ARM AND HE BROKE OUT INTO A RASH. PATIENT THEN BECAME SHORT OF BREATH. EMS WAS CALLED AND PATIENT WAS TRANSPORTED TO HOSPITAL, WHERE HE WAS TREATED FOR ANAPHYLACTIC SHOCK TO THE COVID MODERNA VACCINE.

VAERS ID: 923766 (history)
Form: Version 2.0
Age: 26.0
Sex: Female
Location: New York
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / UNK	RA / -

Administered by: Private **Purchased by:** ?
Symptoms: Gastric disorder, Headache, Laboratory test, Lip swelling, Nausea, Swelling face, Urticaria
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ADDERALL
Current Illness:
Preexisting Conditions: ADHD, Lyme disease.
Allergies: Latex. benzol peroxide
Diagnostic Lab Data: They did preform labs while I was in the hospital but I am not sure what they were.
CDC Split Type: vsafe

Write-up: Went to the ER on the 31st, my face swelled and chest was covered in hives. Lips swelled. I would get nauseated and I felt like there was acid in my stomach. I also had a headache. I have been taking Zyrtec, Benadryl as needed.

VAERS ID: 923822 (history)
Form: Version 2.0
Age: 36.0

Sex: Female
Location: Massachusetts
Vaccinated: 2020-12-21
Onset: 2020-12-22
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Altered state of consciousness, Chills, Diarrhoea, Dizziness, Fatigue, Headache, Hyperhidrosis, Pain, Pneumonia, Productive cough, Pyrexia, Vomiting
SMQs:, Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: The day after the vaccine I had fatigue and body aches. Then on 12-23 fevers of 101.4, chills, body aches, diarrhea, vomiting, productive cough, and profuse sweating. Jan 1st I was in the shower and was dizzy with a headache. Loss of consciousness almost happened and that is when i got out of the shower and laid down. Went to the ED and was diagnosed with bilateral pneumonia

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VAERS ID: 923892 (history)
Form: Version 2.0

Age: 49.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / SYR

Administered by: Work **Purchased by:** ?
Symptoms: Blood pressure increased, Blood test, Dizziness, Electrocardiogram, Fear, Feeling abnormal, Feeling hot, Flushing, Headache, Immediate post-injection reaction, Injection site reaction, Limb discomfort, Pain in extremity, Palpitations, Peripheral coldness, Tremor, Urine analysis
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Parkinson-like events (broad), Noninfectious encephalopathy/delirium (broad), Hypertension (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypersensitivity (narrow), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Ziac (blood pressure) Zoloft Levothyroxine (thyroid medication) Multi Vitamin Calcium Trisprintec (birth control) Generic Sudafed
Current Illness:
Preexisting Conditions: Hypertension Throid
Allergies: Arithimycin Tetracycline
Diagnostic Lab Data: EKG Blood work Urine test
CDC Split Type: vsafe

Write-up: I did let the nurse know right after I got the vaccine It did feel like a huge water balloon was sitting on my arm and very heavy at the site. I went to the waiting area and the last 5 minutes and I started feeling weird, kind of getting dizzy and my heart started racing really fast. Am I can tell I was getting really hot and my face was flushed and my heart was racing and my hands started shaking, trembling almost like I couldn"t control it. I"ve never experienced something like this. The part that really scared me was my heart racing like it was going to come out of my chest. They called a code, put me in a wheel chair and took me to the ER (where I work). They took my blood pressure which was really high, they did EKG, blood work and urine test. They kept me there for about five hours, while watching me. And

the last weird thing that happened at one point I felt like liquid was running thru my body and my feet were really cold for several hours, It felt like someone was pouring liquid over me. I kept asking the nurse if this is normal and no one knew what to tell me. After everything started slowing back down and it had been a couple hours and the next day i felt like I had a slight head ache and felt out of it, kind of like I had a hangover and of course my arm was really sore. The Doctor in the ER did advise me not to get the second dose as it will be worse than the first one.

VAERS ID: 923901 (history)
Form: Version 2.0
Age: 54.0
Sex: Female
Location: Wisconsin
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Dyspnoea, Feeling hot, Flushing
SMQs:, Anaphylactic reaction (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ALBUTEROL INHALER AS NEEDED, ATORVASTATIN 20MG DAILY, CITALOPRAM 20MG DAILY, GLIMEPIRIDE 1 MG DAILY, LISINOPRIL 20MG DAILY. ASPIRIN 81MG DAILY
Current Illness:
Preexisting Conditions: DIABETES, HYPERTENSION, DEPRESSION, ASTHMA
Allergies: PENCILLIN (HIVES, GI UPSET), IODINE (HIVES, RASH), SULFA (UNKNOWN), IODINATED CONTRAST DYE (HIVES), LATEX (HIVES)
Diagnostic Lab Data:
CDC Split Type:

Write-up: LEFT SIDED CHEST PAIN, SHORTNESS OF BREATH, FELT WARM AND FLUSHED

VAERS ID: 923935 (history)
Form: Version 2.0
Age: 62.0
Sex: Female
Location: Massachusetts
Vaccinated: 2020-12-19
Onset: 2020-12-19
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Chills, Cytokine storm, Fatigue, Malaise, Nausea, Oxygen saturation abnormal, Pyrexia, Respiratory failure, SARS-CoV-2 test positive, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (narrow), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (broad), Tumour lysis syndrome (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Hypokalaemia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 11 days
Extended hospital stay? No
Previous Vaccinations: fever and chills post shingles shot
Other Medications: tylenol for arthritis
Current Illness: none
Preexisting Conditions: arthritis
Allergies: morphine, erythromycin, IV iron
Diagnostic Lab Data: rapid covid - positive labs - all normal now
CDC Split Type: vsafe

Write-up: I got my shot on the 19th and that evening it was like a light switch and I was so tired I went to sleep at 730pm I had severe chills and fever and had to go to bed. The next day I still wasn''t feeling well and I was called in to get covid tested and I went to the ER on the 21st and took a rapid covid test that was positive. I was stable and had good oxygenation and was discharged. I have fever nausea vomiting I also had problems with O2 stat i was in the 80s and realized I was having respiratory failure so I was admitted on the 27th and I''ve been here ever since. I had kinetic storm and infusions my O2 stats were bad and I was sent to the covid unit and put on high flow oxygen and negative for a PE, I''m still on the covid unit but I feel much better today

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VAERS ID: 924029 (history)
Form: Version 2.0
Age: 53.0
Sex: Female
Location: New York
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	012620A / UNK	- / IM

Administered by: Public **Purchased by:** ?
Symptoms: Dizziness, Dysphagia, Heart rate increased, Throat tightness
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Vestibular disorders (broad), Hypersensitivity (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: plaquinel synthroid
Current Illness: Lupus Hypothyroidism
Preexisting Conditions:
Allergies: none
Diagnostic Lab Data: Benadryl 50 mg P.O. - I was taken via ambulance to the hospital where I received IV steroids & hydration.

CDC Split Type: N/A

Write-up: after 20-30 minutes my throat started to get tight, I could not swallow properly, I felt dizzy & my heart was beating fast.

VAERS ID: 924050 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: Hawaii
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Dyspnoea
SMQs:, Anaphylactic reaction (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: Zolpidem, chocolate
Diagnostic Lab Data:
CDC Split Type:

Write-up: anaphylaxis, dyspnea

VAERS ID: 924173 (history)
Form: Version 2.0
Age: 41.0
Sex: Male

Location: Michigan
Vaccinated: 2020-12-23
Onset: 2020-12-25
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Catheterisation cardiac, Chest pain, Electrocardiogram, Palpitations, Troponin
SMQs:, Arrhythmia related investigations, signs and symptoms (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: metoprolol succinate XL 25 mg 24 hr tablet Commonly known as: Toprol XL Learn more Take 1 tablet (25 mg total) by mouth daily. cholecalciferol 125 mcg (5,000 unit) tablet Commonly known as: Vitamin D3 ferrous sulfate 325 mg (65 mg iron)
Current Illness: None
Preexisting Conditions: Coronary Artery Disease, Asthma
Allergies: None
Diagnostic Lab Data: I had cardiac troponin , ECG and a cardiac catheterization to rule out major adverse cardiovascular event. I am not sure if the vaccine caused the symptoms, but I strongly suspect it.
CDC Split Type:

Write-up: I started having intermittent chest pain moderate in intensity and palpitations.

VAERS ID: 924188 (history)
Form: Version 2.0
Age: 38.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00

Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Computerised tomogram, Migraine, Nausea, Paraesthesia, Swelling face
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (narrow), Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: no
Current Illness: no
Preexisting Conditions: no
Allergies: no
Diagnostic Lab Data: CT Scan at Boswell Medical Center 10401 W. Thunderbird, Sun City, AZ
CDC Split Type:

Write-up: Migraines, right side of face swollen, nausea, tingling

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VAERS ID: 924201 (history)
Form: Version 2.0
Age: 94.0
Sex: Male
Location: Maryland
Vaccinated: 2020-12-24
Onset: 2021-01-03
Days after vaccination: 10
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Acute coronary syndrome, Back pain, Catheterisation cardiac abnormal, Chest pain, Coronary arterial stent insertion, Coronary artery stenosis, Echocardiogram abnormal, Ejection fraction decreased, Electrocardiogram abnormal, Percutaneous coronary

intervention, Troponin increased
SMQs:, Cardiac failure (narrow), Myocardial infarction (narrow), Arrhythmia related investigations, signs and symptoms (broad), Retroperitoneal fibrosis (broad), Embolic and thrombotic events, arterial (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (narrow), Other ischaemic heart disease (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Prilosec 20 mg qd Lipitor 20 mg qd Levothyroxine 100 mcg/d (incr from 88 for TSH 4.78 8/26/20) Ativan 0.5-1 mg hs prn 10-15/month EC ASA 2 x 81 mg qd
Current Illness: Adjustment disorder/grief reaction
Preexisting Conditions: PAST MEDICAL HISTORY Surgical- T&A age 5 Appendectomy age 8 or 9 Rhinoplasty for deviated septum x 2 1960s Gas gangrene Right Leg- debridement 1972 Right inguinal hernia ~2002 Bilateral Cataracts/IOLs 2014 Umbilical Herniorrhaphy 2020 Medical- Hepatitis and possible malaria 1944 Hyperlipidemia GERD/Barretts- Lower GI bleed hospitalized ~1994; no Tx Hemorrhoids Anal fistula Herpes zoster Nov 2012 Left Trigeminal Hypothyroidism Dec 2013 (fatigue, elevated TSH 5.81, fT4 0.7) Childhood Diseases- Measles- yes; Mumps- yes; Rubella- ?; Varicella- yes; Rheumatic Fever- no; Pertussis- yes Habits- Tobacco- quit 1950; Alcohol- 0-2 night; Drugs- no Immunizations- Td- 3/08, 8/22/19; TdaP- ; Influenza- 8/25/20; Pneumovax- yes <2003, 3/08; PCV13- 12/19/14; Hep A- ; Hep B- ; Zostavax- 3/07; Shingrix- 2/21/18, 10/30/18; Covid 19 (Moderna) 12/24/20
Allergies: PCN- rash? Bee sting- swelling (has Epipen) (no hx food or latex allergy)
Diagnostic Lab Data: Elevated Troponin, NSST changes on ECG, Cardiac Catheterization showing 99% mid-LAD and 30-40% RCA lesions. Echocardiogram +hypokinesis of anterior wall/septum with reduced EF 45-50%
CDC Split Type:

Write-up: Patient tolerated the vaccine well with no apparent side effects. Ten days later awoke 12:30 AM with severe chest and upper back pain, presented to Med Center where he was found to have an Acute Coronary Syndrome. Transferred to Medical Center where he underwent successful PCI with two drug eluting stents for a 99% mid-LAD stenosis

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VAERS ID:	924208 (history)
Form:	Version 2.0
Age:	58.0
Sex:	Male
Location:	Illinois
Vaccinated:	2020-12-30
Onset:	2020-12-31
Days after vaccination:	1
Submitted:	0000-00-00

Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, Fatigue, Pyrexia
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Shortness of breath, fever, fatigue

VAERS ID: 924297 (history)
Form: Version 2.0
Age: 71.0
Sex: Male
Location: New York
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood glucose normal, COVID-19, Unresponsive to stimuli, Urinary tract infection

SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad),

Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Tylenol Tablet 325 MG (Acetaminophen) Aspirin EC Tablet Delayed Release 81 MG (Aspirin) Topamax Tablet 100 MG (Topiramate) Keppra Tablet 750 MG (LevETIRAcetam) Melatonin Tablet 10 MG Multi-Day Plus Iron Tablet (Multiple Vitamins-Iron) Refre
Current Illness: RIGHT LEG POSITIVE (+) EXAM FOR DVT
Preexisting Conditions: CHRONIC OBSTRUCTIVE PULMONARY DISEASE, UNSPECIFIED OTHER GENDER IDENTITY DISORDERS MORBID (SEVERE) OBESITY DUE TO EXCESS CALORIES OTHER SEIZURES INSOMNIA DUE TO OTHER MENTAL DISORDER ESSENTIAL (PRIMARY) HYPERTENSION UNSPECIFIED ATRIAL FIBRILLATION ATHSCL HEART DISEASE OF NATIVE CORONARY ARTERY W/O ANG PCTRS SCHIZOPHRENIA, UNSPECIFIED BENIGN PROSTATIC HYPERPLASIA WITHOUT LOWER URINRY TRACT SYMP TYPE 2 DIABETES MELLITUS WITHOUT COMPLICATIONS
Allergies: No known allergies.
Diagnostic Lab Data: Transferred to hospital; diagnosed with COVID-19 and UTI.
CDC Split Type:

Write-up: At around 11:40am resident was observed to be unresponsive. resident noted with pulse and respiration. Not in any distress. lung sounds clear. Vital Signs BP162/82 P86 R18 T97.1 O2 Sat 96%, fingerstick is 133mg/dl .Resident received COVID 19 vaccine at 11:25am. O2 via 2l NC initiated. Nurse Stat call, 911 initiated, MD at Bedside. Resident awake and responsive. EMT responded and resident left with EMT to be transferred to hospital, remains awake and not in any respiratory distress.

VAERS ID:	924410 (history)
Form:	Version 2.0
Age:	76.0
Sex:	Male
Location:	South Dakota
Vaccinated:	2020-12-28
Onset:	2021-01-05
Days after vaccination:	8
Submitted:	0000-00-00
Entered:	2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Adult failure to thrive, Asthenia, Respiratory failure
SMQs:, Anaphylactic reaction (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (narrow), Guillain-Barre syndrome (broad), Hypersensitivity (broad), Respiratory failure (narrow), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: AMLODIPINE BESYLATE 5MG TAB ASPIRIN 81MG EC TAB LEVOTHYROXINE (SYNTHROID) 88MCG TAB TAMSULOSIN HCL 0.4MG CAP FLUTICASONE 500MCG/SALMETEROL 50MCG INHL DISK 60 ROSUVASTATIN CA 20MG TAB CHOLECALCIFEROL 25MCG (1,000UNIT) TIOTROPIUM 2.5MCG
Current Illness:
Preexisting Conditions: Hypertension, COPD with oxygen dependence, hypothyroidism, osteoporosis
Allergies: None
Diagnostic Lab Data:
CDC Split Type:

Write-up: Adult failure to thrive; Chronic hypoxemic respiratory failure; Generalized weakness

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VAERS ID: 924524 (history)
Form: Version 2.0
Age: 31.0
Sex: Female
Location: Minnesota
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	LA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Hypoaesthesia, Obstructive airways disorder, Pruritus, Throat irritation
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Asthma/bronchospasm (broad),

Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Eosinophilic pneumonia (broad), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Sexual dysfunction (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations: ANAPHYLAXIS TO INFLUENZA VACCINE, UNKNOWN DATE
Other Medications: UNKNOWN
Current Illness: UNKNOWN
Preexisting Conditions: UNKNOWN
Allergies: ANAPHYLAXIS TO INFLUENZA VACCINES
Diagnostic Lab Data: UNKNOWN
CDC Split Type:

Write-up: PATIENT REPORTING ITCHING AT 30 MINUTES POST INJECTION. AT 1.5 HOURS POST INJECTION PATIENT REPORTED ITCHY THROAT AND NUMBESS OF LEFT SIDE OF FACE. AT THAT TIME ADVISED TO GO TO EMERGENCY ROOM. NEXT DAY WHEN I FOLLOWED UP WITH PATIENT, SHE REPORTED HER AIRWAY STARTED TO CLOSE AND SHE RECEIVED EPINEPHRINE, AFTER 5 HOURS HER STARTED TO CLOSE AGAIN AND RECEIVED ANOTHER DOSE OF EPINEPHERINE, WAS RELEASED FROM HOSPITAL ROUGHLY 15-16 HOURS AFTER GOING TO ER.

VAERS ID: 924539 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: California
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / UNK	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Blood test, Dizziness, Flushing, Nausea, Oropharyngeal pain, Retching, Throat tightness
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures

(narrow), Vestibular disorders (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Albuterol Inhaler, Nortriptyline, baclofen
Current Illness: none
Preexisting Conditions: osteio arthritis, joint hyper mobility, chronic migrans, fibromyalgia, IBS,
Allergies: Penicillin and mushrooms, and nsaid.
Diagnostic Lab Data: 01/06/21-Blood panels and vital signs.
CDC Split Type:

Write-up: At 10:12 am, Client c/o of sore throat, tightness in throat that relieve quickly, nausea, dry heaves, flushed, light headed and dizziness. Called for EMT. They arrive at 10:25 am and transported her to the local hospital for observation. 01/06/21-Treatment in hospital blood draw, medications given Zofran, Decadron, Benadryl, and Pepcid, and IV fluids. Discharged home at 1244.

VAERS ID: 924658 (history)
Form: Version 2.0
Age: 92.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-26
Onset: 2021-01-05
Days after vaccination: 10
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Erythema, Hypotension, Oxygen saturation decreased, Sensitive skin, Skin warm, Unresponsive to stimuli
SMQs:, Anaphylactic reaction (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Acute central respiratory depression (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective

pneumonia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Caltrate, Vitamin B, Azopt, Ensure, Sertraline, Dulcolax, Benzonatate, Tylenol, Melatonin, Flonase, Imodium, Ativan.
Current Illness: L98.9 Disorder of the skin and subcutaneous tissue, unspecified-Scalp(Primary, Admission), R41.841 Cognitive communication deficit, R13.10 Dysphagia, unspecified, R13.12 Dysphagia, oropharyngeal phase, C44.311 Basal cell carcinoma of skin of nose, R26.81 Unsteadiness on feet, M62.81 Muscle weakness (generalized), R53.1 Weakness, Z74.09 Other reduced mobility, Z74.1 Need for assistance with personal care, D51.9 Vitamin B12 deficiency anemia, unspecified, D64.9 Anemia, unspecified, E87.6 Hypokalemia, F03.90 Unspecified dementia without behavioral disturbance, F32.9 Major depressive disorder, single episode, unspecified, G47.00 Insomnia, unspecified, H40.9 Unspecified glaucoma
Preexisting Conditions: L98.9 Disorder of the skin and subcutaneous tissue, unspecified-Scalp(Primary, Admission), R41.841 Cognitive communication deficit, R13.10 Dysphagia, unspecified, R13.12 Dysphagia, oropharyngeal phase, C44.311 Basal cell carcinoma of skin of nose, R26.81 Unsteadiness on feet, M62.81 Muscle weakness (generalized), R53.1 Weakness, Z74.09 Other reduced mobility, Z74.1 Need for assistance with personal care, D51.9 Vitamin B12 deficiency anemia, unspecified, D64.9 Anemia, unspecified, E87.6 Hypokalemia, F03.90 Unspecified dementia without behavioral disturbance, F32.9 Major depressive disorder, single episode, unspecified, G47.00 Insomnia, unspecified, H40.9 Unspecified glaucoma
Allergies: PCN, Sulfa
Diagnostic Lab Data: All assessment and laboratory data is being managed by the hospital,not this facility at this time, the resident is still admitted to the hospital and they areattempting to stabilize
CDC Split Type:

Write-up: Severe Hypotension, Redness, Warmth and sensitivity all over skin surfaces, lack of responsiveness, low oxygen saturation.

VAERS ID:	924722 (history)
Form:	Version 2.0
Age:	96.0
Sex:	Female
Location:	Illinois
Vaccinated:	2021-01-02
Onset:	2021-01-06
Days after vaccination:	4

Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0142 / UNK	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Tylenol, amlodipine, ASA, Atorvastatin, combivent, cyclobensap
Current Illness: neck fracture, CKD 3, HTN, HL, edema, heart disease
Preexisting Conditions: asthma
Allergies: cortisone, morphine
Diagnostic Lab Data: None
CDC Split Type:

Write-up: Severe shortness of breath, administered inhaler, hydralazine with no improvement. Dr. notified. Sent to ER

VAERS ID: 927830 (history)
Form: Version 2.0
Age: 71.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / -

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Bilirubin urine present, Chills, Chromaturia, Laboratory test, Liver function test increased, Nausea, Vomiting

SMQs:, Rhabdomyolysis/myopathy (broad), Liver related investigations, signs and symptoms (narrow), Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Biliary system related investigations, signs and symptoms (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Synthroid, Tagamet, Lipitor, Protonix
Current Illness: None
Preexisting Conditions: Hypthyroid, GERD, Arthritis
Allergies: NKA
Diagnostic Lab Data: ongoing hospitalization with multiple tests
CDC Split Type:

Write-up: Abdominal pain, chills, n/v, dark urine, elevated LFT"s, Bilirubin in urine. Patient currently admitted to hospital

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VAERS ID: 927938 (history)
Form: Version 2.0
Age: 73.0
Sex: Male
Location: Hawaii
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	- / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Chest X-ray abnormal, Pneumonia, Pyrexia, SARS-CoV-2 test positive
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Received Moderna vaccine on 12/29/2020. 12/30/2020 fever of 100.4 Tylenol given and monitored and fever went down. 12/31/2020 Chest x-ray completed and sent to the hospital and admitted with pneumonia. 1/3/2021 reported by the hospital that Covid-19 results were positive. He had had Covid-19 positive results back on 11/4/2020 prior to vaccine.

VAERS ID: 922339 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-16
Onset: 2020-12-17
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553-L450 / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Heart rate, Heart rate increased, Throat tightness, Tremor
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Noninfectious encephalopathy/delirium (broad), Hypersensitivity (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days

Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201217; Test Name: Pulse rate; Result Unstructured Data: Test Result:178/83
CDC Split Type: GBPFIZER INC2020517453

Write-up: Cramp to throat and body trembling rapid heart beat, ambulance called and taken to hospital Pulse rate 178/83; Cramp to throat and body trembling rapid heart beat, ambulance called and taken to hospital Pulse rate 178/83; Cramp to throat and body trembling rapid heart beat, ambulance called and taken to hospital Pulse rate 178/83; This is a spontaneous report from a contactable consumer downloaded from the Medicines Agency (MA) Regulatory Authority-WEB. Regulatory authority report number is GB-MHRA-ADR 24544791. An 81-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Lot number EJ0553-L450), parenteral on 16Dec2020 as single dose for covid-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient experienced cramp to throat and body trembling rapid heart beat, ambulance called and taken to hospital pulse rate 178/83 on 17Dec2020, which were serious per hospitalization. The patient underwent lab tests and procedures which included heart rate: 178/83 on 17Dec2020. The outcome of cramp to throat and body trembling rapid heart beat, ambulance called and taken to hospital pulse rate 178/83 was recovered on 17Dec2020. No follow-up attempts are possible. No further information is expected.

VAERS ID: 923152 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Body temperature, Cough, Hypertension, Malaise, Polymerase chain reaction, Pyrexia
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Hypertension (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201228; Test Name: blood pressure; Result Unstructured Data: Test Result:200 to 100 mmHg; Test Date: 20201228; Test Name: temperature; Result Unstructured Data: Test Result:below 38 Centigrade; Comments: below 38 C; Test Name: PCR test; Result Unstructured Data: Test Result:no result
CDC Split Type: DEPFIZER INC2020517238

Write-up: Blood pressure 200 to 100 mm Hg; cough; slight symptoms like after an influenza vaccine; temperature below 38 C; This is a spontaneous report from a contactable physician. A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number: unknown), via an unspecified route of administration on 28Dec2020 at single dose (dose 1) for COVID-19 immunization. The patient medical history and concomitant medications were not reported. The patient had blood pressure 200 to 100 mm Hg, temperature below 38 C, cough, slight symptoms like after an influenza vaccine on 28Dec2020. The reaction started on 28Dec2020 (during the night). The patient was hospitalized on an unspecified date in Dec2020. No results from PCR test (unspecified date). Measures taken consisted of "monitoring". Event outcome was recovering. Lot/batch number has been requested.; Sender''s Comments: The reported events temperature below 38 C, cough and slight symptoms like after an influenza vaccine were likely related to the single dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), while the blood pressure increase with high value of 200 to 100 mmHg was unlikely causally related to the BNT162B2. The case will be reassessed should additional information become available, especially patient''s age and underlying diseases. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory authorities, Ethics committees and Investigators, as appropriate.

VAERS ID: 923153 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-27
Onset: 2020-12-27
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site /

		Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	E-J6796 / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Accidental overdose, Asthenia, Fatigue, Headache, Malaise, Pain
SMQs:, Drug abuse and dependence (broad), Guillain-Barre syndrome (broad), Medication errors (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: DEPFIZER INC2020519333

Write-up: malaise; fatigue; body aches; 8 vaccinees have been injected with five times the dose; Headache; weakness; The initial case was missing the following minimum criteria: unspecified number of patients. Upon receipt of follow-up information on 28Dec2020, this case now contains all required information to be considered valid. This is a spontaneous report from a contactable Physician. This Physician reported similar events for eight patients. This is the fourth of eight reports. A 49 years female patient received the 1st dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot E-J6796, expiry date: Apr2021) via Intramuscular on 27Dec2020 at single dose in the upper arm for COVID-19 immunisation. Medical history and concomitant medications were not reported. The patient presented herself at the hospital emergency room after consultation with the public health department and the hospital. The patient experienced Headache, general weakness, malaise. The patient did not wish to be admitted as an inpatient yesterday. A basic diagnosis was initiated via the hospital. The patient is now hospitalized as of this afternoon. On 27Dec2020, employee of a care home received the five-times dose of the corona vaccine supplied by BioNtech/Pfizer. They have also clarified that the vials administered were diluted so a large volume was administered. The patient is still hospitalized with headache, weakness and fatigue but is stable now. information received from BioNTech, which included "To add further colour to the picture, the 8 overdoses happened at one singular care home with one vaccinator only - these were not separate episodes. He is an 82 years old paediatrician who has avidly vaccinated before. The vaccine was diluted, but of course he did not remember that it contained 5 individual doses." Physician said that enclosed is the current status from 29Dec2020, 2 p.m. After consultation with the vaccinator and the vaccination nurse, the vaccine was reconstituted by the nurse (with 1.8 ml NaCl) and drawn into a 2 ml syringe until it stopped (2.25 ml). This total amount was given to all 8 vaccinees. The patient experienced headache and weakness on vaccination day, on 28Dec2020 Headache at times, body aches, increasing weakness. Announced presentation in the emergency room, remains under inpatient observation. Outcome of "8 vaccinees have been injected with five times the dose" was unknown, while outcome of other events was not

recovered. This case is being treated according to the Act. No life-threatening adverse event(s), ambulatory treatment and Hospitalization required.; Sender"s Comments: Based on the time association, the events are possibly related to suspect BNT162b2 injection with overdose. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.,Linked Report(s) : DE-PFIZER INC-2020515377 Same reporter, same product, different patient

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VAERS ID: 923159 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-16
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-06

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Body temperature, Headache, Lethargy, Photophobia, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (narrow), Glaucoma (broad), Corneal disorders (broad), Retinal disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ;
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201216; Test Name: Body temperature; Result Unstructured Data: Test Result:40 Centigrade

CDC Split Type: GBPFIZER INC2020517955

Write-up: Headache and fever/fever of 40; Headache, fever, lethargy; Photophobia; Headache, fever, lethargy; This is a spontaneous report from a contactable physician downloaded from the Medicines Agency (MA) WEB GB-MHRA-EYC 00235575 , Sender"s (Case) Safety Report Unique Identifier : GB-MHRA-ADR 24544051. A 56-years-old female patient received her dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) (Lot Number: EJ0553), Intramuscular on 15Dec2020 at single dose for COVID-19 vaccination. The patient medical history was not reported. Concomitant medication included ramipril, atorvastatin, amlodipine and carbamazepine. The patient experienced headache and fever, lethargy, photophobia on 16Dec2020. One day after receiving vaccine developed fever of 40 with headache and lethargy and mild photophobia on 16Dec2020. The events were reported as serious with the criteria of hospitalization. The outcome of the events was unknown. No follow-up attempts are possible. No further information is expected.

VAERS ID: 924820 (history)
Form: Version 2.0
Age: 81.0
Sex: Male
Location: Colorado
Vaccinated: 2020-12-31
Onset: 2021-01-02
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chills, Cough, Dysuria, Micturition urgency, Nausea, Pneumonia, Pollakiuria, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: PENICILLIN (UNKNOWN, REMOTE REACTION)

Diagnostic Lab Data:
CDC Split Type:

Write-up: COUGH, RIGORS, NAUSEA, VOMITING, URINARY URGENCY/FREQUENCY, DYSURIA - FOUND TO HAVE LLL PNEUMONIA, CONCERNING FOR POSSIBLE CYRPTOGENIC ORGANIZING PNEUMONIA

VAERS ID: 924822 (history)
Form: Version 2.0
Age: 82.0
Sex: Female
Location: Unknown
Vaccinated: 2020-12-29
Onset: 2020-12-31
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: ATIVAN METOPROLOL
Diagnostic Lab Data:
CDC Split Type:

Write-up: PATIENT DEVELOPED PROGRESSIVE NEW DYSPNEA, DIFFERENT FROM HER BASELINE. SHE HAS BEEN HOSPITALIZED TWICE FOR PERSISTENT DYSPNEA AND CENTRALIZED CHEST PAIN, WHICH HAS OTHERWISE HAD NEGATIVE WORK UP.

VAERS ID: 924835 (history)

Form: Version 2.0
Age: 23.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-01-04
Onset: 2021-01-05
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025520A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood test normal, Chest X-ray normal, Chills, Dizziness, Electrocardiogram normal, Headache, Injection site pain, Nausea, Pain, Pyrexia, Tachycardia, Troponin normal, Urine analysis normal
SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: Diagnosed with covid December 1st, recovered 3 and a half weeks prior to vaccine
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: Troponin - normal Blood Panel - normal Chest x-ray- normal Urine analysis- normal EKG- normal showed tachycardia
CDC Split Type:

Write-up: 103.5 Fever that wouldn?t come down with Tylenol, chills, sharp headache, tachycardia, site pain, dizziness, body aches, nausea All symptoms started 11 hours after first dose of vaccine (3AM), went to hospital 15 hours after symptoms started and was treated for 9 hours until all symptoms abruptly stopped

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VAERS ID: 925078 (history)
Form: Version 2.0
Age: 39.0

Sex: Male
Location: New York
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0140 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Cardiac arrest, Hypotension, Oxygen saturation decreased
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Cardiomyopathy (broad), Respiratory failure (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 10 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Reglan 5mg, Maalox Advance 20mg, Prednisone 2.5 mg, Metoprolol succinate ER 25mg, Prednisone Lisinopril 2.5 mg, Eliquis 5mg, Atorvastatin 80mg, Gabapentin300mg, Tylenol 325mg x2, multivitamin, iron 325mg, magnesium oxide 400mg, zinc 220mg,
Current Illness: N/A
Preexisting Conditions: acute cerebral vascular insufficiency, aphasia, cerebral infarction, gastro reflux, vitamin deficiency, long use systemic steroid, central pain syndrome, dermatitis, cardiomyopathy, crones disease, SOB
Allergies: N/A
Diagnostic Lab Data: Inpatient
CDC Split Type:

Write-up: 12/29/2020 2 hr after vaccination patient became hypotensive, decreased oxygen levels was transferred to Hospital currently inpatient at hospital - admitted for cardiac arrest

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VAERS ID: 925254 (history)
Form: Version 2.0
Age: 40.0

Sex: Female
Location: Texas
Vaccinated: 2020-12-22
Onset: 2020-12-23
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / SYR

Administered by: Work **Purchased by:** ?
Symptoms: Chills, Fatigue, Full blood count normal, Heart rate irregular, Influenza like illness, Mobility decreased, Pain in extremity, Pyrexia, SARS-CoV-2 test negative, Septic screen
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Parkinson-like events (broad), Cardiac arrhythmia terms, nonspecific (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Levaquin
Diagnostic Lab Data: Covid test Sepsis test full panel blood work
CDC Split Type: vsafe

Write-up: I woke the next morning with flu like symptoms my arm was hurting, I was feelin tired and really couldn"t get out of bed. As the day progressed I got chills, started running a fever, It went up to 104 and my heart rate went up and down from 120-165 so I went to the ER. They gave me fluids and ran a whole bunch of test, tested me for Covid and Sepsis and everything came back normal and negative for Covid. They monitored my heart rate and once it started getting back to normal, they ended up letting me go home, I was there from 8pm to about 3am. After that I felt tired and felt like when I had Covid back in November. I started feeling better about Sunday, still a little tired but felt more back to normal. My heart rate is still getting back to normal so my Dr is following me on that.

VAERS ID: 925756 (history)
Form: Version 2.0

Age: 66.0
Sex: Female
Location: Louisiana
Vaccinated: 2020-12-29
Onset: 2020-12-31
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / -

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram, Echocardiogram, Hypoaesthesia, Hypoaesthesia oral, Laboratory test, Magnetic resonance imaging, Pain, SARS-CoV-2 test negative
SMQs:, Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), COVID-19 (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; BENTYL; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Hypertension; Low back pain; Reflux gastritis
Allergies:
Diagnostic Lab Data: Test Name: Cat scan; Result Unstructured Data: Test Result:Unknown results; Test Name: ultrasound of heart; Result Unstructured Data: Test Result:Unknown results; Test Name: labs; Result Unstructured Data: Test Result:Unknown results; Test Name: MRI; Result Unstructured Data: Test Result:Unknown results; Test Date: 20210101; Test Name: Nasal Swab/ Covid 19; Result Unstructured Data: Test Result:Negative
CDC Split Type: USPFIZER INC2021003473

Write-up: Right sided facial & top lip Numbness & recurring pain; Right sided facial & top lip Numbness & recurring pain; Right sided facial & top lip Numbness & recurring pain; This is a spontaneous report from a contactable consumer reporting for herself. A 66-year-old female patient (not pregnant at the time of vaccination) received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number EK9231), via unspecified route of administration on 29Dec2020 13:30 on left arm at single dose for COVID-19 immunization. Facility where the most recent COVID-19 vaccine was administered in Hospital. Medical history included Hypertension, Reflux and low back pain. Concomitant medications included ibuprofen, dicycloverine hydrochloride (BENTYL), Valsartan and omeprazole. The patient previously took codeine, Bactrim, Zyrtec and experienced allergies. The patient experienced

Right sided facial and top lip Numbness, recurring pain on 31Dec2020 16:00. All events resulted in emergency room visit and Hospitalization on 31Dec2020 for 1 day. The patient underwent lab tests and procedures, which included Cat scan, MRI, ultrasound of heart and labs on unspecified dates, Nasal Swab/Covid 19 test on 01Jan2021 with negative result. The outcome of the events were not resolved.

VAERS ID: 925777 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Michigan
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anion gap, Blood calcium decreased, Blood chloride increased, Blood creatinine normal, Blood glucose increased, Blood potassium normal, Blood sodium normal, Blood urea normal, Carbon dioxide decreased, Eye swelling, Haematocrit normal, Haemoglobin normal, Hypoaesthesia, Mean cell haemoglobin concentration normal, Mean cell haemoglobin normal, Mean cell volume normal, Pain in jaw, Platelet count normal, Red blood cell count normal, Red cell distribution width normal, SARS-CoV-2 test negative, Swelling face, Swollen tongue, Throat tightness, White blood cell count increased
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (narrow), Angioedema (narrow), Peripheral neuropathy (broad), Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Osteonecrosis (broad), Chronic kidney disease (broad), Hypersensitivity (narrow), Tumour lysis syndrome (broad), Tubulointerstitial diseases (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None

Preexisting Conditions: None, just tree nut allergy with history of anaphylaxis to tree nut last episode 2011.
Allergies: Tree Nuts
Diagnostic Lab Data: Lytes: Na 137, K 4.8, Cl 107, CO2 21, Anion Gap 9, BUN 12, Creatinine 0.85, Calcium 8.6, glucose 223, WBC 11, RBC 4.41, hemoglobin 13.4, hematocrit 40.4, MCV 91.6, MCH 30.4, MCHC 33.2, RDW 13.3, Platelet Count 335, COVID PCR negative
CDC Split Type:

Write-up: Felt sharp pain under jaw, then facial numbness. Then quickly developed facial swelling, left eye swelling and tongue swelling. Then felt like throat closing up within 7 minutes of receiving vaccine. Received vaccine in the clinic, was transported to Emergency Room. In ER received Epinephrine, steroids, Benadryl and Pepcid.

VAERS ID: 925925 (history)
Form: Version 2.0
Age: 52.0
Sex: Male
Location: New York
Vaccinated: 2020-12-28
Onset: 2021-01-04
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	LA / -

Administered by: Work **Purchased by:** ?
Symptoms: Computerised tomogram, Electroencephalogram, Magnetic resonance imaging, Pyrexia, Seizure
SMQs:, Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Leviteracetam for seizures because he had a first time seizure after testing positive for COVID-19. Gabapentin Osteo Bio Flex Vitamin D, Vitamin C
Current Illness: None
Preexisting Conditions: None

Allergies: None
Diagnostic Lab Data: EEG , MRI, CAT SCAN
CDC Split Type:

Write-up: 1 day after the vaccine he had a low grade fever... 1 week later he had a seizure and then multiple ones in the ER...

VAERS ID: 926042 (history)
Form: Version 2.0
Age: 41.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Cough, Dyspnoea, Swollen tongue
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Famotidine, Protonix
Current Illness:
Preexisting Conditions: GERD
Allergies: Allergy to sulfa, latex, bee-stings
Diagnostic Lab Data:
CDC Split Type:

Write-up: Developed shortness of breath, swelling of tongue, persistent cough within 5 minutes of vaccination. Was treated with EpiPen and kept in ER for observation overnight.

Symptoms resolved.

VAERS ID: 926221 (history)
Form: Version 2.0
Age: 64.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-18
Onset: 2020-12-19
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood immunoglobulin G, Cough, Diarrhoea, Fatigue, Full blood count, Liver function test, Metabolic function test, SARS-CoV-2 antibody test, SARS-CoV-2 test positive, T-lymphocyte count
SMQs:, Anaphylactic reaction (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 8 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Levothyroxine 50mcg daily , Aloglitpan Benzoate 25mg daily, alendronate 70mg once weekly.
Current Illness: 11/20/20 appointment for mid-thoracic back pain.
Preexisting Conditions: Essential Hypertension, Hyperlipidemia, Hypothyroid, Type 2 diabetes with renal complication, without insulin , Chronic renal disease., BMI 36.58
Allergies: NKA
Diagnostic Lab Data: Daily Hepatic function, Basic Metabolic Panel, complete blood count. 1/5: Immunoglobulin G, Covid 19 IGG Antibody, T-Cell Subset panel, T4/T8 T-Cell subset panel
CDC Split Type:

Write-up: Patient contacted provider 12/26 with following symptoms: Dry cough, diarrhea, fatigue for 7 days. Covid test ordered and positive. presented to ED on 12/31 and admitted into hospital. Still inpatient as of 1/7/2021.

VAERS ID: 926230 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Echocardiogram, Electrocardiogram, Full blood count, Laboratory test, Sinus tachycardia, Supraventricular tachycardia
SMQs:, Supraventricular tachyarrhythmias (narrow), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Symbicort Levothyroxine Myrbetriq vitamin D vitamin C Calcium
Current Illness:
Preexisting Conditions: Asthma Hashimotos Thyroiditis Overactive bladder
Allergies: Penicillin Clindamycin
Diagnostic Lab Data: EKG CBC Chemistry ECHO
CDC Split Type:

Write-up: Patient experienced an episode of SVT and then sinus tachycardia for approximately 6 hours after injection

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VAERS ID: 926433 (history)
Form: Version 2.0
Age: 88.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-28
Onset: 2020-12-31
Days after vaccination: 3

Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram abnormal, Haemorrhagic stroke, Intensive care, Neurologic neglect syndrome, Visual impairment
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Anticholinergic syndrome (broad), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Glaucoma (broad), Optic nerve disorders (broad), Lens disorders (broad), Retinal disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Elequis, Amiodoron, pravastatin, losartin, synthroid, timalol
Current Illness: high bp, high cholesterol,
Preexisting Conditions: deaf in rt ear - rt vestibular schwanoma,
Allergies: none
Diagnostic Lab Data: 2 CT scans showing large bleed. ,
CDC Split Type:

Write-up: Hemmoragic Stroke. Began with vision difficulty in the morning. Then I noticed she had left sided neglect. Went to ER. Treated with Andresxa (to counteract Elaquis). In SICU for 2 nights then telemetry unit for 3 nights. CUrrently in Rehab.

VAERS ID: 926454 (history)
Form: Version 2.0
Age: 62.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-04
Onset: 2021-01-05
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal adhesions, Abdominal pain, Surgery

SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Hyperlipidemia DJD
Allergies: Avelox, Dilaudid, Percocet, Pylera
Diagnostic Lab Data: Surgery 1/6/2021
CDC Split Type:

Write-up: Following vaccination the patient had progressively worsening abdominal pain over the next 24 hours. Presented to the ER and was initially thought to have appendicitis. However, it was then discovered during surgery that the appendix was surgically absent. The surgeon did not that the patient did have a "Round, infarcted ligamentous tissue was wrapped around ascending colon. "

VAERS ID: 926462 (history)
Form: Version 2.0
Age: 91.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-28
Onset: 2021-01-04
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1042 / 1	UN / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Absence of immediate treatment response, Death, Hypoxia
SMQs:, Asthma/bronchospasm (broad), Lack of efficacy/effect (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Infective pneumonia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 2021-01-05
Days after onset: 1
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Lorazepam, pantoprazole, miralax, senna plus, carbodopa-levodopa, metoprolol, mirtazipine, quetiapine
Current Illness: dementia, Upper gastrointestinal bleed
Preexisting Conditions: Advanced dementia, Parkinson"s" disease ,chronic kidney disease, sciatica, hyperlipidemia
Allergies: no known food or medication allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient developed hypoxia on 1/4/2021 and did not respond to maximal treatment and passed way on 1/5/2021

VAERS ID: 926568 (history)
Form: Version 2.0
Age: 77.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-28
Onset: 2020-12-30
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1042 / 1	UN / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Absence of immediate treatment response, Death
SMQs:, Lack of efficacy/effect (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-04
Days after onset: 5
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: alprazolam, amlodipine, ASA, cilostazol, advair, lamotrigine, ,losartan, metoprolol,omprazole,simvastatin
Current Illness: chronic medical problems as noted below
Preexisting Conditions: dementia, peripheral vascular disease, COPD, hypertension, GERD, S/P CVA with stent of left carotid ,right common iliac and right external iliac arteries, CDK,diabetes
Allergies: clopidigrel, sulfa, vicodin
Diagnostic Lab Data:
CDC Split Type:

Write-up: patient declined 12/30/2020 and was transferred to hospital where he did not respond to treatment and passed away 1/4/2020

VAERS ID: 926703 (history)
Form: Version 2.0
Age: 35.0
Sex: Male
Location: Oklahoma
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J202A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Blood test, Guillain-Barre syndrome, Immunoglobulin therapy, Magnetic resonance imaging, Neuralgia, Neurological examination, Paraesthesia
SMQs:, Peripheral neuropathy (narrow), Guillain-Barre syndrome (narrow), Demyelination (narrow), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Bradycardia at rest. History of Guillain Barre Syndrome/AIDP in 2010.
Allergies: None

Diagnostic Lab Data: MRI and bloodwork, neuro screening and tests.
CDC Split Type:

Write-up: Guillain Barre syndrome/AIDP event. Paresthesia and nerve pain developed in bilateral legs 4 hours after shot and progressed slowly for 4 days in intensity and area involved. Symptoms progressed distally to superior. On the 5th day symptoms progressed rapidly and involved bilateral legs up to the groin, left arm up to lateral shoulder, and right hand. I went to the hospital and was admitted to start IVIG treatment for Guillain Barre Syndrome/AIDP.

VAERS ID: 926755 (history)
Form: Version 2.0
Age: 70.0
Sex: Female
Location: New Mexico
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram, Electroencephalogram, Foaming at mouth, Incoherent, Laboratory test, Magnetic resonance imaging, Respiratory arrest, Resuscitation, Seizure, Speech disorder, Unresponsive to stimuli
SMQs:, Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Dementia (broad), Convulsions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (narrow), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Hypersensitivity (broad), Respiratory failure (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: warfarin, lisinopril, atenolol, Lasix, potassium, Tylenol
Current Illness: rheumatic heart disease, hx of stroke, brain hemorrhage, open heart X3 with valve replacement, VP shunt
Preexisting Conditions: rheumatic heart disease, hx of stroke, brain hemorrhage, open

heart X3 with valve replacement, VP shunt
Allergies: codeine
Diagnostic Lab Data: CT , MRI and labs done, eeg conducted due to her previous medical HX.
CDC Split Type:

Write-up: Vaccine Candidate received vaccine approxat 2:30pm, was monitored for 15 min no complications at the time, went home. Around 5:30pm while walking into her home she became unresponsive, was assisted in a siting position, became incoherent, mumbling and started to convulse to the right side of her r upper extremity. Foaming at the mouth and stopped breathing, CPR was initiated for 1-2 min, EMS arrived was transported to Medical Center. She was admitted and is currently hospitalized. MD reports this event is highly unlikely related to the vaccine given her medical history but suggested to report being its a new vaccine. Current status: stable.

VAERS ID: 926778 (history)
Form: Version 2.0
Age: 63.0
Sex: Male
Location: New York
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	012L20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Oedema peripheral, Pyrexia, Unresponsive to stimuli
SMQs:, Cardiac failure (broad), Angioedema (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypotonic-hyporesponsive episode (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: none known, person did
Preexisting Conditions: Developmentally disabled, history of lower extremity edema

Allergies: None known
Diagnostic Lab Data: N/A
CDC Split Type:

Write-up: Person had a fever of 102.4, pulse rate of 118, he was non-responsive with edema of the right calf & ankle this morning when he was assessed.

VAERS ID: 926787 (history)
Form: Version 2.0
Age: 53.0
Sex: Female
Location: Minnesota
Vaccinated: 2020-12-30
Onset: 2020-12-31
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / UNK	- / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Asthenia, Blood glucose normal, Breath sounds abnormal, Confusional state, Consciousness fluctuating, Depressed level of consciousness, Dyspnoea, Fatigue, Heart rate increased, Lethargy, Muscle twitching, Oxygen saturation decreased, Pyrexia, Seizure like phenomena, Tremor
SMQs:, Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (narrow), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Convulsions (narrow), Dyskinesia (broad), Dystonia (broad), Parkinson-like events (broad), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: gabapentin, asa, oxycodone, fentanyl, flexiril, requip, omeprazole, keflex, symbicort, restasis, spiriva, synthroid, tylenol, simvastatin, lasix, aldactone
Current Illness:
Preexisting Conditions: DM2, COPD, chronic uti/ neurogenic bladder, CKD, obesity,

depression,
Allergies: bee stings, abilify, flowers,
Diagnostic Lab Data: Completed in the ED hospital and then transferred.
CDC Split Type:

Write-up: Resident had the COVID vaccine 12/30/2020. 12/31/20, resident has been in bed all shift. Staff became concerned when resident was not easily aroused. Resident displayed signs of tremors, twitching, confusion, in and out of consciousness, low O2 sats, elevated pulse and fever, fatigue and weakness. Writer called NP. NP stated this is most likely a reaction d/t the COVID vaccine. She gave orders for Benadryl 25mg IM x1 now and Tylenol 1000 mg now. NP also stated resident will not be getting the second dose of vaccine. Will continue to monitor and update NP if worsening symptoms. After receiving Benadryl and Tylenol at 145pm, resident began to appear as though she was feeling better and was talking to talk, fever had gone down. Tonight resident is not easily aroused, lethargic, continues to have tremors and twitches, almost appearing as convulsions. When asked if she knows where she is or what day it is, resident can properly answer. Resident denies SOB but staff has noted loud squeals while breathing. NP was updated and gave new orders to give Benadryl 25 mg IM x1 if needed and Ok to send resident to ED. Resident currently refuses to go to the hospital. Will continue to monitor. BP 152/112, P 116, T 99.1, O2 87-91. Resident''s O2 at 1205am was 80% on 3LPM. Resident unable to be aroused from sleep by writer. NAR called to assist. NAR could not arouse resident. Writer and NAR attempted to reposition resident and resident''s breathing became more labored. Resident turned back to previous position and writer called on call MD at approx. 1220am. MD returned call approx. 1235am with orders to send resident to ED. 911 called and ambulance arrived about 1245am. History of present condition given to EMTs and they stated resident would be going to Hospital. Writer has attempted to contact Hospital ED x3 but have been unable to get through. An EMT did just call to clarify when vaccine was given, what symptoms have been present and when they started. She said she has everything she should need and she will let Hospital ED staff know to call if they need anything else. Writer will again attempt to contact them though. Resident''s temp was 97.5 and BG 128. When EMTs arrived they got an O2 reading of 60%. Resident did open her eyes a couple times during transfer from bed to stretcher and while stretcher was going outside but no responses from resident were made.

VAERS ID: 927006 (history)
Form: Version 2.0
Age: 68.0
Sex: Female
Location: South Carolina
Vaccinated: 2020-12-31
Onset: 2021-01-01
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Arthropathy, C-reactive protein increased

SMQs:, Arthritis (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: clopidogrel, carvedilol, atorvastatin, aspirin, albuterol, torsemide, buproprion
Current Illness: None
Preexisting Conditions: HFrEF, CAD, HTN, COPD, depression, OSA, ckd
Allergies: latex, nickel
Diagnostic Lab Data: CRP $g300
CDC Split Type:

Write-up: Diffuse polyarthropathy starting the day after vaccination and continuing for 7+ days. Currently treating with abx for concern of possible cellulitis, and prednisone 60mg for polyarthropathy. Currently admitted.

VAERS ID: 927065 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Mississippi
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EA9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest X-ray abnormal, Computerised tomogram normal, Laboratory test abnormal, Oropharyngeal pain, Pneumonia, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Flecinide 100 mg/day
Current Illness: none
Preexisting Conditions: vitriol fibulation
Allergies: none
Diagnostic Lab Data: Pneumonia - chest x-ray-29th test for it CT scan - negative
CDC Split Type: vsafe

Write-up: woke up with fever and sore throat on 22nd; went to job and got tested and tested positive; on the 23rd developed right upper lobe pneumonia; on 27th was hospitalized with three lobe pneumonia; On 22nd received got zpack - azithromycin and sudafed; received at hospital doxicycleln IV ; ivermectin and went home with them, as well. Hospital

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VAERS ID: 927096 (history)
Form: Version 2.0
Age: 60.0
Sex: Male
Location: Massachusetts
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood test, Chest X-ray, Chills, Computerised tomogram abdomen, Computerised tomogram head, Computerised tomogram thorax abnormal, Echocardiogram, Fatigue, Flushing, Generalised tonic-clonic seizure, Hyperhidrosis, Legionella test, Magnetic resonance imaging brain, Myalgia, Pulmonary embolism, Pyrexia, SARS-CoV-2 test, Scan with contrast, Streptococcus test, Ultrasound abdomen
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Convulsions (narrow), Embolic and thrombotic events, venous (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Generalised convulsive seizures following immunisation (narrow), Hypersensitivity (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Venlafaxine 150mg daily Ibuprofen 600mg 2x/week
Current Illness: None
Preexisting Conditions: Mild osteoarthritis
Allergies: N/A
Diagnostic Lab Data: CT head without contrast Blood work CT abdomen/pelvis with contrast Xray chest Covid-19 RT-PCR Legionella and Strep Urine Echo TTE MRI brain with and without contrast MRA head without contrast US abdomen CT chest Pulmonary embolism with contrast
CDC Split Type:

Write-up: Day 2 (12/29/20): Fever (<100 degrees), Mild muscle aches, Fatigue Day 3 (12/30/20): Fatigue, Muscle aches Day 4 (12/31/20): Alternating chills and profuse sweating starting at 8am, Full body flushing, Grand Mal Seizure at 4:30pm

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VAERS ID: 927260 (history)
Form: Version 2.0
Age: 87.0
Sex: Female
Location: Wisconsin
Vaccinated: 2020-12-28
Onset: 2021-01-06
Days after vaccination: 9
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	AR / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood lactic acid increased, Bradycardia, Cardiac arrest, Computerised tomogram head normal, Death, Heart sounds abnormal, Pulse absent, Respiration abnormal, Resuscitation, SARS-CoV-2 test negative, Unresponsive to stimuli, White blood cell count increased
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Lactic acidosis (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 2021-01-06
Days after onset: 0
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vit D3, Miralax, Spironolactone, Tramadol, Metoprolol Succinate, MOM, Ferrous Gluconate, Ondansetron, Nitroglycerin Sublingual PRN, PreserVision AREDS, Fluticasone Propionate Suspension, Lisinopril, Clopidogrel Bisulfate, Campor-Methol-Met
Current Illness: Digestive Surgery for bowel obstruction
Preexisting Conditions: HTN, AAA, Chronic Diastolic Heart Failure, Diabetes Type 2, Paroxysmal atrial fibrillation, bradycardia, Hx of STEMI, Hx of CVA, HX of uterine cancer,
Allergies: Codeine, Meperidine, Morphine, Estrogens, Penicillins, Tetanus Toxoids
Diagnostic Lab Data: Brain CT negative WBC and lactic acid increased. COVID 19 on 1/4/21 negative
CDC Split Type:

Write-up: No adverse effects noted after vaccination. Patient with cardiac history was found unresponsive at 16:45 on 1/6/21. Abnormal breathing patterns, eyes partially closed SPO2 was 41%, pulseless with no cardiac sounds upon auscultation. CPR and pulse was regained and patient was breathing. Patient sent to Hospital ER were she remained in an unstable condition had multiple cardiac arrest and severe bradycardia and in the end the hospital was unable to bring her back.

VAERS ID: 937030 (history)
Form: Version 2.0
Age: 58.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-23
Onset: 2020-12-23
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Hypoaesthesia, Hypoaesthesia oral, Laboratory test normal
SMQs:, Peripheral neuropathy (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Sexual dysfunction (broad)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: LEFT FACIAL AND TONGUE NUMBNESS - WORK UP CVA NEG Narrative: Developed left facial and tongue numbness 4 hours after vaccine - went to ER and admitted for 2 days, negative workup for CVA or other acute etiology. Symptoms resolved prior to discharge from hospital

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VAERS ID: 940544 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: Unknown
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure increased, Heart rate increased, Hypertension, Nausea, Pharyngeal swelling, Tachycardia, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypertension (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: NauseaVomiting, HYPERtension, Tachycardia, throat swelling Narrative: Went home after vaccine and starting vomiting, throat swelling. HR and BP elevated Went to ER was given beta blockers, in hospital for observation.

VAERS ID: 950443 (history)
Form: Version 2.0
Age: 43.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-28
Onset: 2020-12-30
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Angina pectoris, Malaise, Myalgia, Tachycardia, Tachypnoea
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Eosinophilic pneumonia (broad), Other ischaemic heart disease (narrow), Respiratory failure (broad), Tendinopathies and ligament disorders (broad), Infective pneumonia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Tachypnea, Angina, Tachycardia, Sore Muscles Narrative: Started feeling ill on the 28th. Drove himself to get checked out at the hospital. Hospital sent him home. Called EMS at 12/30/2020 @ 0200 and was hospitalized for 1 day. He is more stable now but still has these symptoms at a moderate level.

VAERS ID: 925613 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-28
Onset: 2020-12-01
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Dyspnoea, Inflammatory marker test, Respiratory tract infection, SARS-CoV-2 test, X-ray
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Infective pneumonia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Type 2 diabetes mellitus
Preexisting Conditions: Medical History/Concurrent Conditions: Cardiac valve disease (underwent surgery)
Allergies:
Diagnostic Lab Data: Test Name: Inflammatory markers; Test Result: Positive ; Test Name: COVID-19 virus test; Test Result: Negative ; Comments: 3 times; Test Name: X-ray; Result Unstructured Data: Test Result:compatible with respiratory infection; Comments: positive, possible pneumonia

CDC Split Type: GRPFIZER INC2021001570

Write-up: respiratory tract infection; breathing difficulty; This is a spontaneous report from a contactable physician (Pfizer employee). A 54-year-old male patient (hospital physician) received BNT162B2 (COMIRNATY) via intramuscular at single dose on 28Dec2020 for COVID-19 immunisation. Medical history included ongoing diabetes type 2; valve disease for which he had undergone a surgery. Concomitant medications were not reported. The patient experienced breathing difficulty 1-2 days after vaccination with respiratory tract infection in Dec2020. And he was hospitalized since Dec2020. He was intubated at the ICU possibly about 1 day later and was transferred to the ICU of another hospital. Patient"s X-ray was compatible with respiratory infection (positive, possible pneumonia), inflammatory markers were positive, while COVID-19 virus test was negative 3 times. The outcome of events was not recovered. The information on the lot/batch number has been requested.; Sender"s Comments: The Company cannot completely exclude the possible causality between the reported respiratory infection (positive, possible pneumonia) with breathing difficulty and the administration of BNT162B2, based on the reasonable temporal association. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RA, IEC, as appropriate.

VAERS ID: 925614 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-27
Onset: 2020-12-30
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Asthenia, Dizziness, Headache, Malaise, Myalgia, Vision blurred
SMQs:, Rhabdomyolysis/myopathy (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Glaucoma (broad), Lens disorders (broad), Eosinophilic pneumonia (broad), Retinal disorders (broad), Vestibular disorders (broad), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: ILPFIZER INC2020521523

Write-up: does not feel well.; blurred vision; dizziness; Strong headache; muscle pains; general weakness; This is a spontaneous report from a contactable consumer via a consumer (patient"s father). A male patient of an unspecified age received bnt162b2 (lot/batch number and expiration date not provided), via an unspecified route of administration, on 27Dec2020, at single dose, for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. It was reported that the patient was a vaccinated this past Sunday (27Dec2020) for COVID-19. He is in hospital and does not feel well. Dizziness, blurred vision, strong headache, muscle pains, general weakness. These events started just today (30Dec2020). The outcome of the event was unknown. Information about batch number has been requested.

VAERS ID: 925621 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Abdominal pain
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: PTPFIZER INC2020521543

Write-up: Severe abdominal pain, cramp like; This is a spontaneous report from a contactable physician. This report was received via a sales representative. A 32-years-old female patient received bnt162b2 (COMIRNATY) via an unspecified route of administration on an unspecified date at 0.3 mL, single dose for covid-19 immunisation. Medical history and concomitant medications were not reported. The patient, a nurse, experienced severe abdominal pain, cramp like, 12 h after administration of the Covid-19 vaccine. The patient remained at the hospital overnight in observation. The patient was treated with the following medicaiton for the events: buthylescapolamine, ondasetron, paracetamol and 2mg. The patient was discharged the next day, in the morning. At the time of the report, the outcome of the event was unknown. Information on batch number has been requested.; Sender''s Comments: Based on the compatible temporal association, a possible contributory role of vaccination with bnt162b2 (COMIRNATY) in the development of abdominal pain cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 925622 (history)
Form: Version 2.0
Age: 87.0
Sex: Male
Location: Foreign
Vaccinated: 2020-12-26
Onset: 2020-12-26
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK4241 / UNK	LA / OT

Administered by: Other **Purchased by:** ?
Symptoms: Activated partial thromboplastin time, Alanine aminotransferase, Amylase, Aspartate aminotransferase, Blood bilirubin, Blood chloride, Blood creatinine, Blood glucose, Blood potassium, Blood sodium, Blood urea, C-reactive protein, Gamma-glutamyltransferase, Haematocrit, Haemoglobin, Investigation, Pain in extremity, Platelet count, Protein urine, Prothrombin time, Red blood cell count, Vaccination site haematoma, White blood cell count
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Proteinuria (narrow), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No

Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: DUROFILIN; HEFEROL [FERROUS FUMARATE]; PRESOLOL [LABETALOL HYDROCHLORIDE]; LASIX [FUROSEMIDE]; PRILENAP [ENALAPRIL MALEATE;HYDROCHLOROTHIAZIDE]; FOLACIN [FOLIC ACID]
Current Illness: Hypertension
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201229; Test Name: aPTT; Result Unstructured Data: Test Result:1.13 R; Test Date: 20201229; Test Name: ALT; Result Unstructured Data: Test Result:14; Test Date: 20201229; Test Name: Amylase; Result Unstructured Data: Test Result:102; Test Date: 20201229; Test Name: AST; Result Unstructured Data: Test Result:17; Test Date: 20201229; Test Name: Billirubin; Result Unstructured Data: Test Result:4.4/1.3; Test Date: 20201229; Test Name: Chloride; Result Unstructured Data: Test Result:106; Test Date: 20201229; Test Name: Creatinine; Result Unstructured Data: Test Result:145; Test Date: 20201229; Test Name: SUK; Result Unstructured Data: Test Result:5.3; Test Date: 20201229; Test Name: Potassium; Result Unstructured Data: Test Result:5.4; Test Date: 20201229; Test Name: Natrium; Result Unstructured Data: Test Result:141; Test Date: 20201229; Test Name: urea; Result Unstructured Data: Test Result:16.4; Test Date: 20201229; Test Name: C-reactive protein; Result Unstructured Data: Test Result:2.4; Test Date: 20201229; Test Name: GGT; Result Unstructured Data: Test Result:15; Test Date: 20201229; Test Name: Haematocrit; Result Unstructured Data: Test Result:037; Test Date: 20201229; Test Name: Haemoglobin; Result Unstructured Data: Test Result:119; Test Date: 20201229; Test Name: Local finding; Result Unstructured Data: Test Result:Normal vascular status,; Comments: palpable peripheral pulse, no signs of venous insufficiency; Test Date: 20201229; Test Name: Platelet; Result Unstructured Data: Test Result:150; Test Date: 20201229; Test Name: UP; Result Unstructured Data: Test Result:72; Test Date: 20201229; Test Name: PT; Result Unstructured Data: Test Result:0.97 R; Test Date: 20201229; Test Name: Red blood; Result Unstructured Data: Test Result:4.1; Test Date: 20201229; Test Name: Leucocytes; Result Unstructured Data: Test Result:3.93
CDC Split Type: RSPFIZER INC2020521535

Write-up: hematoma on the lateral side of the left upper arm; on the first day he had pain in his left arm; This is a spontaneous report from a contactable physician received from the Agency of Regulatory Authority. The regulatory authority report number is 515-00-01723-2020-2. An 87-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number: EK4241; Expiration date: Apr2021), intramuscularly in the left arm, on 26Dec2020 (at the age of 87-year-old) as a single dose for COVID-19 immunisation. Medical history included ongoing hypertension. Concomitant medications included theophylline (DUROFILIN), ferrous fumarate (HEFEROL), labetalol hydrochloride (PRESOLOL), furosemide (LASIX), enalapril maleate, hydrochlorothiazide (PRILENAP), and folic acid (FOLACIN)). The patient previously received influenza vaccine (MANUFACTURER UNKNOWN) (the patient has been receiving a influenza vaccine for several years and has never experienced adverse reactions or bigger hematomas at the injection site) on an unspecified date for immunization and experienced no adverse event. The patient experienced hematoma on the lateral side of the left upper arm on 26Dec2020, which caused hospitalization from 29Dec2020 to an unspecified date. The patient also experienced on the

first day he had pain in his left arm on 26Dec2020. The clinical course was reported as follows: Before the vaccine was administered, the epidemiologist who conducted immunization oversight noticed multiple bruises on the arm, as well as a bruise on the right leg of the patient. The patient reported that the bruises were appearing and disappearing for a long time. The attending physician stated that there were no coagulation disorders in the medical history, nor other hematology disorders; as for the chronic illnesses, only hypertension. The patient was not taking anticoagulant therapy. The patient has been receiving a flu vaccine for several years and has never experienced adverse reactions or bigger hematomas at the injection site. It was reported that when administering each dose of the vaccine, aspiration was done, and the blood vessels were checked for any punctures. On 26Dec2020 in the elderly nursing home, the patient received the vaccine against COVID-19. After several hours, they noticed a hematoma in the area of the left brachium. On the first day, the patient felt pain in the left arm. He did not have fever. The patient was admitted on 29Dec2020 for observation. At admission, the patient negated the arm pain and stated that hematoma had not expanded. It was reported that the hematoma on the lateral side of the left brachium is most probably a result of accidental damage to superficial blood vessel by a needle during the administration of the vaccine. The arm was without pain, swelling, redness or warmth at the injection site, and patient was afebrile. The general condition was good, blood pressure within normal values, as well as his pulse. The patient underwent lab tests and procedures which included, all on 29Dec2020: activated partial thromboplastin time (aPTT): 1.13 R, alanine aminotransferase (ALT): 14, amylase: 102, aspartate aminotransferase (AST): 17, bilirubin: 4.4/1.3, chloride: 106, creatinine: 145, SUK: 5.3, potassium: 5.4, natrium: 141, urea: 16.4, C-reactive protein: 2.4, gamma-glutamyltransferase (GGT): 15, haematocrit: 037, haemoglobin: 119, local finding: normal vascular status, palpable peripheral pulse, no signs of venous insufficiency, platelet: 150, urine protein (UP): 72, prothrombin time (PT): 0.97 R, red blood: 4.1, and leucocytes: 3.93. The clinical outcome hematoma on the lateral side of the left upper arm was unknown and of on the first day he had pain in his left arm was recovered on an unspecified date. It was also reported that at the time when the information was obtained, the reporter did not know whether the patient was still hospitalized and whether the hematoma was diminishing or not.

VAERS ID: 925679 (history)
Form: Version 2.0
Age: 33.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood count, Feeling hot, Heart rate, Malaise, Medical observation, Paraesthesia oral, Physical examination, Tachycardia, Tryptase, Vein disorder
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions

(excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Blood count; Result Unstructured Data: Test Result:unknown results; Test Date: 20201217; Test Name: Heart rate; Result Unstructured Data: Test Result:120; Test Name: Medical observation; Result Unstructured Data: Test Result:Normal; Test Date: 20201217; Test Name: Physical examination; Result Unstructured Data: Test Result:Normal; Test Date: 20201217; Test Name: Blood tryptase; Result Unstructured Data: Test Result:unknown results; Test Date: 20201217; Test Name: Blood tryptase; Result Unstructured Data: Test Result:unknown results
CDC Split Type: GBPFIZER INC2021001743

Write-up: lips tingling; Felt hot; veins bulging; tachycardic, heart rate 120; feel unwell; This is a spontaneous report from a contactable physician and other healthcare professional from the Medicines Agency (MA) regulatory authority-WEB GB-MHRA-ADR 24545598. A 33-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 17Dec2020 at single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. On 17Dec2020, patient left the observation area slightly early but started to feel unwell and came straight back. Felt hot, lips tingling, veins bulging, tachycardic, heart rate 120. Not hypotensive. Not breathless. Assessed by general practitioner and paramedic. Intramuscular adrenaline, intramuscular chlorphenamine and intramuscular hydrocortisone given. Ambulance called. Felt hadn''t significantly improved. Additional dose adrenaline may have been given prior to ambulance arrival. Oxygen administered. Intramuscular adrenaline given by ambulance and admitted to emergency department. Patient felt well in herself. Her observations were normal. She had no signs of adverse reaction now. Physical exam was normal. Time was currently 6 hours post event. She had been reviewed by emergency department consultant. She had been advised to attend general practitioner/ minor injuries unit next day for a 24-hour post event tryptase. She was happy with the plan. The patient underwent lab tests and procedures which included blood count with unknown results, and medical observation: normal both on unknown date, heart rate: 120, physical examination: normal, blood tryptase twice with unknown results all on 17Dec2020. Outcome of all events was recovered on 17Dec2020. No follow-up attempts possible. No further information expected, information about lot/batch number cannot be obtained.

VAERS ID: 925684 (history)
Form: Version 2.0

Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-14
Onset: 2020-12-15
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Facial paresis, Hemiparesis, Magnetic resonance imaging
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Magnetic resonance imaging; Result Unstructured Data: Test Result:t2 signal flare in periventriculr region
CDC Split Type: GBPFIZER INC2021004496

Write-up: Weakness left arm, face and leg; Facial weakness; This is a spontaneous report from a contactable consumer downloaded from the Agency Regulatory Authority-WEB GB-MHRA-EYC 00235898, GB-MHRA-ADR 24545298. A 34-years-old female patient received bnt162b2 (COMIRNATY), parenteral on 14Dec2020 at single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient previously took isotretinoin (ROACCUTANE) in 2017 and experienced hemiplegic migraine. The patient experienced weakness left arm, face and leg on 15Dec2020. Seek advice details included speaking to general practitioner. Referred to hospital. Magnetic resonance imaging scan showed t2 signal flare in periventriculr region. The events outcome was not recovered. No follow-up attempts are possible, information about batch number cannot be obtained.

VAERS ID: 925689 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood glucose, Blood pressure decreased, Blood pressure measurement, Electrocardiogram, Heart rate, Heart rate decreased, Muscle twitching, Oxygen saturation, Presyncope, Respiratory rate, Unresponsive to stimuli
SMQs:, Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dyskinesia (broad), Dystonia (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Aneurysm repair (Abdominal aneurysm repair); Myocardial infarction (2 x previous MI)
Allergies:
Diagnostic Lab Data: Test Date: 20201217; Test Name: Blood glucose; Result Unstructured Data: Test Result:8.6; Test Date: 20201217; Test Name: Blood pressure; Result Unstructured Data: Test Result:dropped; Test Date: 20201217; Test Name: Blood pressure; Result Unstructured Data: Test Result:130/60; Test Date: 20201217; Test Name: Blood pressure; Result Unstructured Data: Test Result:116/57; Test Date: 20201217; Test Name: Blood pressure; Result Unstructured Data: Test Result:120/60; Test Date: 20201217; Test Name: Electrocardiogram; Result Unstructured Data: Test Result:unknown results; Test Date: 20201217; Test Name: Pulse rate; Result Unstructured Data: Test Result:decreased; Test Date: 20201217; Test Name: Pulse rate; Result Unstructured Data: Test Result:50; Test Date:

20201217; Test Name: Pulse rate; Result Unstructured Data: Test Result:59; Test Date: 20201217; Test Name: Pulse rate; Result Unstructured Data: Test Result:69; Test Date: 20201217; Test Name: Oxygen saturation; Test Result: 98 %; Test Date: 20201217; Test Name: Oxygen saturation; Test Result: 98 %; Test Date: 20201217; Test Name: Oxygen saturation; Test Result: 98 %; Test Date: 20201217; Test Name: Respiratory rate; Result Unstructured Data: Test Result:12; Test Date: 20201217; Test Name: Respiratory rate; Result Unstructured Data: Test Result:16; Test Date: 20201217; Test Name: Respiratory rate; Result Unstructured Data: Test Result:12
CDC Split Type: GBPFIZER INC2021002179

Write-up: Twitching; Unresponsive to stimuli; Pulse decreased; Blood pressure dropped; Vasovagal attack; This is a spontaneous report from a contactable other healthcare professional downloaded from the Medicines Agency (EMA) EudraVigilance-WEB GB-MHRA-WEBCOVID-20201217165432, Safety Report Unique Identifier GB-MHRA-ADR 24544494. A 98-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) Batch/lot number: EJ0553, via an unspecified route of administration on 17Dec2020 at single dose for COVID-19 vaccination. Medical history included myocardial infarction (2 x previous MI), aneurysm repair (Abdominal aneurysm repair) both unknown if ongoing. The patient"s concomitant medications were not reported. The patient experienced vasovagal attack on 17Dec2020, twitching on 17Dec2020 15:17, unresponsive to stimuli on 17Dec2020, pulse decreased on 17Dec2020, blood pressure dropped on 17Dec2020. The events were serious as hospitalization. Clinical course: Patient was given vaccine and had almost completed the 15 minute observation when his son noticed he was "twitching" and not responding. This happened at 15:17. Patient was transferred to a wheelchair and moved to the resus area where he was guided onto the trolley. He was became more lucid at this stage. A full set of observations was taken (blood pressure 130/60, pulse 69, oxygen saturation 98%, respiratory rate 16, blood sugar 8.6, unable to get a temperature) patient was responding at this stage. Staff member arrived to help also. Patient was responsive for a good 20 minutes but then had another "twitching" episode. Blood pressure dropped to 116/57, pulse down to 50, oxygen saturation 98%, respiratory rate 12. IV (intravenous) access gained at 5.40 and 500ml normal saline commenced. Ambulance called as patient not recovering from vasovagal attack at this point. Ambulance crew arrived shortly afterwards. ECG (electrocardiogram) taken, lying and standing blood pressure also. Further observations taken - blood pressure 120/60, pulse 59, respiratory rate 12, oxygen saturation 98%. Patient helped to stand by ambulance crew and experienced another "twitching" episode so crew took to ED (emergency department) for further tests. Patient was in resus for approximately 1 hour. Patient had not tested positive for COVID-19 (coronavirus diseases 2019) since having the vaccine. The outcome of the events was unknown. No follow-up attempts possible. No further information expected.

VAERS ID: 925696 (history)
Form: Version 2.0
Age: 80.0
Sex: Male
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-01
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553/V0001 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Body temperature, Diarrhoea, Feeling hot, Lethargy, Melaena, Pyrexia, SARS-CoV-2 test
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Gastrointestinal haemorrhage (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Gastrointestinal bleed; Hypertension; Peripheral vascular disease
Allergies:
Diagnostic Lab Data: Test Name: Body temperature; Result Unstructured Data: Test Result:37.7 Centigrade; Test Name: COVID-19 virus test; Test Result: Negative ; Comments: No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2021001657

Write-up: Low grade fever/Body temperature 37.7 centigrade; Melaena; Lethargic; Feeling hot; diarrhoea; This is a spontaneous report from a contactable physician downloaded from the Medicines Agency (MA) Regulatory authority-WEB GB-MHRA-WEBCOVID-20201218181357, Safety Report Unique Identifier GB-MHRA-ADR 24544914. An 80-years-old male patient received dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) (Lot# EJ0553/V0001), via an unspecified route of administration on 15Dec2020 at single dose for COVID-19 vaccination. Medical history included gastrointestinal bleed from an unknown date and unknown if ongoing, peripheral vascular disease and hypertension. Concomitant medication included clopidogrel for peripheral vascular disease, amlodipine for hypertension, pravastatin for peripheral vascular disease. The patient previously took clopidogrel and experienced gastrointestinal bleed. The patient experienced feeling hot/felt hot/lethargic that evening on 15Dec2020, day after and since then diarrhoea (Dec2020)/more formed stool but all melaena on 16Dec2020. The patient had no abdominal pain. The patient experienced mild low grade fever, body temperature 37.7 centigrade on an unspecified date. The patient was admitted to hospital due to all events. The events were reported as serious. This may all be secondary to a gastrointestinal bleed from clopidogrel but wanted to highlight reaction in case a pattern emerges. The patient underwent lab tests and procedures which included COVID-19 virus test on unknown date indicated No - negative COVID-19 test. Patient had not had symptoms associated with COVID-19. Patient was not enrolled in clinical trial. The outcome of the event melaena was not resolved and the outcome of the other events was unknown. No

follow-up attempts possible. No further information is expected.

VAERS ID: 925731 (history)
Form: Version 2.0
Age: 51.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1688 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Headache, Hypertension, Paraesthesia
SMQs:, Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Guillain-Barre syndrome (broad), Hypertension (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Asthma; Comments: Unsure if patient has had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial.
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021004060

Write-up: Headache; Hypertension; Pins and needles in right lower leg; This is a spontaneous report from a other healthcare professional downloaded from the Medicines Agency (EMA) EudraVigilance-WEB , Regulatory Authority number: GB-MHRA-ADR 24545844, Safety Report Unique Identifier [GB-MHRA-WEBCOVID-20201221195950]. A 51-years-old female patient received bnt162b2 (COMIRNATY, lot number: EJ1688), via an unspecified route of administration on 21Dec2020 at single dose for covid-19 immunisation. Medical history included asthma from an unknown date and unknown if ongoing. Unsure if patient has had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial. The patient"s

concomitant medications were not reported. The patient experienced headache, hypertension and pins and needles in right lower leg on 21Dec2020. Patient had pins and needles in right lower leg. Occurred with in 15 minutes of vaccinations. As symptoms not resolving taken to accident and emergency. Patient has not tested positive for COVID-19 since having the vaccine. The outcome of the events was not recovered. The seriousness criteria was reported as hospitalization. No follow-up attempts possible. No further information expected.

VAERS ID: 925732 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2021-01-04
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Body temperature, Chest pain, Heart rate, Heart rate increased, Hypertension, Pyrexia, SARS-CoV-2 test
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hypertension (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19 (tested positive for)
Allergies:
Diagnostic Lab Data: Test Name: BP; Result Unstructured Data: Test Result:156/105; Test Name: temp; Result Unstructured Data: Test Result:38.7; Test Name: pulse; Result Unstructured Data: Test Result:130; Test Date: 202011; Test Name: covid-19; Result Unstructured Data: Test Result:POSITIVE

CDC Split Type: GBPFIZER INC2021006890

Write-up: high temperature of 38.7; Chest pain; high pulse of 130; high blood pressure of 156/105; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) Batch/lot no unknown, via an unspecified route of administration on 04Jan2021 at single dose for covid-19 vaccination. Medical history included tested positive for covid-19 on Nov2020. The patient''s concomitant medications were not reported. The patient was in hospital with a high temperature of 38.7 , chest pain, high pulse of 130 and blood pressure of 156/105 on an unspecified date. The outcome of the events was unknown. The information on the batch/lot number and expiration date has been requested.

VAERS ID: 927519 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Minnesota
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Anaphylactic reaction, Intensive care, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2020521806

Write-up: Anaphylactic reaction; hives in the first 10 minutes of the vaccine; This is a spontaneous report from a contactable Other Health Professional (Physician Assistant). A 29-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 29Dec2020 at a single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The reporter stated that she is treating a patient in ICU that got the COVID-19 vaccine on 29Dec2020. The patient developed hives in the first 10 minutes of the vaccine and had an anaphylactic reaction 1 hour later. Seriousness of events was reported to be hospitalization. Outcome of the events was unknown. The reporter also mentioned that it was not a mild reaction and patient was still in the ICU, 48 hours later. Information about lot/batch number has been requested.; Sender''s Comments: A possible causal association between administration of BNT162B2 and the onset of anaphylactic reaction and hives cannot be excluded, considering the plausible temporal relationship and the known adverse event profile of the suspect product. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 927571 (history)
Form: Version 2.0
Age: 21.0
Sex: Female
Location: New York
Vaccinated: 2020-12-31
Onset: 2020-12-31
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / OT

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Ageusia, Anosmia, Anxiety, Asthenia, Chills, Cough, Crying, Dizziness, Dyspnoea, Fatigue, Heart rate increased, Injection site pain, Myalgia, Nausea, Oropharyngeal pain, Pyrexia, Rhinorrhoea, SARS-CoV-2 test positive
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Taste and smell disorders (narrow), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Depression (excl suicide and self injury) (broad), Vestibular disorders (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Dehydration (broad), Opportunistic infections (broad), Immune-mediated/autoimmune disorders (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19; Penicillin allergy
Allergies:
Diagnostic Lab Data: Test Date: 20201231; Test Name: Heart beat; Result Unstructured Data: Test Result:fast; Test Date: 20210101; Test Name: covid; Test Result: Positive
CDC Split Type: USPFIZER INC2021000409

Write-up: tested positive for Covid test; tested positive for Covid test; difficulty breathing; chills; fluctuating fever; nausea; weakness; weakness/extreme fatigue; loss of taste and smell; loss of taste and smell; muscle pain; cough; sore throat; nasal drip; dizziness; fast heartbeat; injection site pain; anxiety; crying; This is a spontaneous report from a contactable healthcare professional. This 21-year-old female patient reported for herself that she received BNT162B2 1st dose on 31Dec2020 10:00 AM intramuscular at left arm for COVID-19 immunisation. Medical history included known allergies: Penicillin and Covid-19. Concomitant therapy included BC as reported. The patient experienced difficulty breathing, chills, fluctuating fever, nausea, dizziness, weakness, fast heartbeat, tiredness, loss of taste and smell, muscle pain, injection site pain, anxiety, cough, sore throat, nasal drip, crying, extreme fatigue, Etc on 31Dec2020 at 06:00 PM. The events resulted in doctor or other healthcare professional office/clinic visit, emergency. The patient was hospitalized for 1 day and received treatment included blood thinner rivaroxaban (XARELTO) and had 2 weeks quarantine. The patient had Covid prior to vaccination and tested positive for Covid test post vaccination on 01Jan2021. The outcome of the events was not resolved. Information on Lot/batch number has been requested.; Sender''s Comments: Based on available information, a possible contributory role of the subject product, BNT162B2 vaccine, cannot be excluded for the suspected LOE, SARS-CoV-2 test positive and the other reported events due to temporal relationship. Of note, it is reported that the patient had history of COVID 19 infection prior to the vaccination. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics, counteractive treatment measures and concomitant medications. This case will be reassessed once additional information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID:	927981 (history)
Form:	Version 2.0
Age:	56.0
Sex:	Female

Location: Georgia
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Computerised tomogram, Dysarthria, Headache, Hypoaesthesia oral, Lip swelling, Magnetic resonance imaging, Nasal discomfort, Paraesthesia oral, Repetitive speech, Urticaria
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Taking Fioricet 50-325-40 MG Tablet, Sig: 1 tablet- 2 tablets as needed for tension headache Orally Once a day Taking Escitalopram Oxalate 10 MG Tablet, Sig: 1 tablet Orally Once a day Start Date: 04/02/2020 Taking Depo-Estradiol 5 MG/M
Current Illness:
Preexisting Conditions: N95.8 Other specified menopausal and perimenopausal disorders J30.2 Other seasonal allergic rhinitis F34.1 Dysthymia N95.1 Hot flashes due to menopause Z79.890 Postmenopausal HRT (hormone replacement therapy) G43.909 Migraine J45.901 Unspecified asthma with (acute) exacerbation J40 Bronchitis R74.8 Elevated liver enzymes R60.9 Edema J31.0 Rhinitis, unspecified type
Allergies: MSG - ANGIODEMA, Celexa, Cipro - rash and trouble breathing, Imitrex - stops her from breathing, Amitriptyline HCl - drowsiness, Concerta, Ritalin, Tamiflu, Topamax - memory loss, Depakote - Bell''s palsy, Gabapentin, Inderal LA - Lack of therapeutic, Cymbalta, sertraline - headaches
Diagnostic Lab Data: transported to ER 01/05/2021 CT scan 01/05/2021 MRI 01/06/2021
CDC Split Type:

Write-up: Pt experiencing and c/o left nasal burning, left upper lip tingling progressing to numbness with slight swelling noted, scattered patchy hives to upper front chest, sharp HA above right eye, denies SOB, no acute respiratory distress noted or reported, slurring of words shortly after onset of other symptoms. Pt repeating "something ain''t right". Pt received Moderna COVID vaccine at 4:35pm with no reactions or side effect noted within the post 15 and 30 minutes. EMS notified at 5:49pm. MD notifed and ordered Benadryl 50mg IM (given at

5:53pm), EpiPen and DepoMedrol 40mg IM if needed. No respiratory distess noted, pt denies SOB. EMS arrived and transported pt to ER.

VAERS ID: 927983 (history)
Form: Version 2.0
Age: 68.0
Sex: Female
Location: Virginia
Vaccinated: 2020-12-30
Onset: 2020-12-31
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Allen's test, Base excess decreased, Blood bicarbonate normal, Blood pH normal, COVID-19 pneumonia, Computerised tomogram thorax abnormal, Dyspnoea, Fatigue, Lung infiltration, PCO2 decreased, PO2 normal, Pneumonia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Lactic acidosis (broad), Interstitial lung disease (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: celexa
Current Illness: none
Preexisting Conditions: anemia
Allergies: codeine
Diagnostic Lab Data: COVID + ARTERIAL BLOOD GAS: (COLL: 01/07/2021 13:15) (MsgRcvd 01/07/2021 13:58) Final results Test Result Flag **(Reference)** O2 METHOD NASAL CANN FiO2% 28 PUNCTURE SITE RIGHT BRACHIA pH 7.429 (7.350 - 7.450) pCO2 34 L mm/HG (35 - 45) pO2 88.7 mm/HG (75.0 - 100) HCO3 22.8 mm/HG (22.0 - 26.0) BASE EXCESS -1.1 mmol/L O2 SAT 97.1 % (91.0 - 99.0) ALLEN TEST N/A Patchy bilateral infiltrates compatible with pneumonia. Differential diagnosis includes COVID pneumonia shown on CTA of chest.
CDC Split Type:

Write-up: Developed SOB and fatigue 1 day after vaccine, went to urgent care and tested positive for COVID at urgent care. Returned to our ED after going to urgent care again on 1/7/21, had o2 sat of 84% on room air, impoved to 98 on 3 liters. Was transferred to another facility for admission.

VAERS ID: 928062 (history)
Form: Version 2.0
Age: 85.0
Sex: Male
Location: Massachusetts
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood lactic acid increased, Cardiac arrest, Death, Endotracheal intubation, Hypotension, Hypoxia, Laboratory test, Lethargy, Liver function test, Lung infiltration, Pancytopenia, Pulmonary congestion, Pulmonary imaging procedure abnormal, Troponin, Vomiting
SMQs:, Torsade de pointes/QT prolongation (broad), Cardiac failure (broad), Anaphylactic reaction (broad), Acute pancreatitis (broad), Agranulocytosis (narrow), Angioedema (broad), Asthma/bronchospasm (broad), Haematopoietic cytopenias affecting more than one type of blood cell (narrow), Lactic acidosis (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Interstitial lung disease (narrow), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Myelodysplastic syndrome (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-07
Days after onset: 2
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: ACETAMINOPHEN CARBIDOPA-LEVODOPA CARBIDOPA-LEVODOPA ER CITALOPRAM HBR DONEPEZIL HCL EUCERIN FINASTERIDE FLOMAX LATANOPROST OMEPRAZOLE QUETIAPINE FUMARATE QUETIAPINE FUMARATE SENNA LAX
Current Illness: Parkinson"s Disease with advanced dementia, dysphagia. Alcoholism in remission. HTN. BPH. GERD
Preexisting Conditions: Parkinson"s Disease with advanced dementia, dysphagia. Alcoholism in remission. HTN. BPH. GERD
Allergies: angioedema due to ACE and ARB meds
Diagnostic Lab Data: Pancytopenia, elevated lactate, troponin, LFTs. Left mid lung infiltrate and pulmonary vascular congestion.
CDC Split Type:

Write-up: vomiting later on 01/05/21. Lethargy and hypoxia in pm of 01/06/21. Hypotension am of 01/07/21. Hospitalized, intubated, cardiac arrest, died 01/07/21.

VAERS ID: 928291 (history)
Form: Version 2.0
Age: 69.0
Sex: Female
Location: Wisconsin
Vaccinated: 2021-01-04
Onset: 2021-01-05
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Abdominal pain upper, Alanine aminotransferase increased, Aspartate aminotransferase increased, Blood alkaline phosphatase increased, Blood bilirubin increased, Chest pain, Hepatic enzyme increased, Liver function test increased
SMQs:, Liver related investigations, signs and symptoms (narrow), Acute pancreatitis (narrow), Biliary system related investigations, signs and symptoms (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: Cevimeline, diphenhydramine, escitalopram, fluticasone, folic acid, hydroxychloroquine, loratadine, multivitamins, pantoprazole, Synthroid. methotrexate and etanercept. These two medications were last taken on 12/28/2020 and then held by
Current Illness:
Preexisting Conditions: hypothyroidism, sjogren"s syndrome, sicca syndrome, celiac sprue
Allergies: gluten and penicillin
Diagnostic Lab Data: 1/8/2021 AST:1890, ALT:1201, Alk Phos: 167, Total Bili:1.8
CDC Split Type:

Write-up: symptoms:chest and stomach pain Has markedly elevated liver function tests that were normal 2 weeks prior to immunization Is being admitted the hospital to monitor liver function test.

VAERS ID: 928333 (history)
Form: Version 2.0
Age: 76.0
Sex: Male
Location: New Mexico
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20-2A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Adjusted calcium increased, Asthenia, Blood calcium increased, Cough, Hypercalcaemia, Pyrexia, Somnolence
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: insulin detemir 10 units daily losartan 25mg daily simvastatin 60mg daily glipizide 5mg daily vitamin D calcium
Current Illness: none known
Preexisting Conditions: sarcoidosis h/o colon cancer in remission
Allergies: lisinopril - swelling

Diagnostic Lab Data: calcium 13 - corrected calcium 14
CDC Split Type:

Write-up: Fever, coughing, drowsiness, generalized weakness. Was found to be hypercalcemic (corrected calcium 14) and admitted 1/2-4. No prior history of hypercalcemia.

VAERS ID: 928378 (history)
Form: Version 2.0
Age: 95.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 2	UN / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, Hospice care, Myocardial infarction, Respiratory tract congestion, Tachycardia, Unresponsive to stimuli
SMQs:, Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Embolic and thrombotic events, arterial (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: On: Norvasc 5mg daily Losartan 50mg daily Menest 0.3mg daily Vitamin C 500mg twice a day
Current Illness: N/A
Preexisting Conditions: HX of CHF Pneumonia HTN Cardiac Murmur Osteoarthritis
Allergies: Vasotec, Zocor, and Cymbalta
Diagnostic Lab Data: Unknown, will need to follow up with hospital.

CDC Split Type:

Write-up: Congestion Shortness of breath Tachycardia Transferred out 911. Per hospital, patient had a myocardial infarction, is unresponsive, and on hospice services.

VAERS ID: 928461 (history)
Form: Version 2.0
Age: 67.0
Sex: Male
Location: Florida
Vaccinated: 2020-12-29
Onset: 2021-01-03
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Computerised tomogram abnormal, Eye swelling, Intensive care, Laboratory test, Lip swelling, Oedema, Pharyngeal oedema, Pruritus, Rash, Rash erythematous, Rash papular, Swelling face
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Angioedema (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Oropharyngeal allergic conditions (narrow), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Metformin, Lisinopril, Simvastatin, Ozempic , pantopazol
Current Illness: none
Preexisting Conditions: Diabetes
Allergies: Cephlasporin
Diagnostic Lab Data: Multiple labs, ct scan soft tissue edema of throat
CDC Split Type:

Write-up: Anaphylactic reaction, Severe edema and raised red rash entire body, Severe itching ,Soft tissue edema of throat. Swelling of, eyes, lips, face. Multiple trips to ER, treated with steroids, Benadryl, prevacid. , CURRENTLY IN ICU ON EPINEPHRINE DRIP, STEROIDS, MULTIPLE MEDS

VAERS ID: 928558 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Iowa
Vaccinated: 2020-12-30
Onset: 2021-01-04
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / UNK	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Cough, Dialysis, Fatigue, Mechanical ventilation, SARS-CoV-2 test positive
SMQs:, Acute renal failure (narrow), Anaphylactic reaction (broad), Acute central respiratory depression (broad), Chronic kidney disease (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Lupus
Allergies:
Diagnostic Lab Data: Unknown--.
CDC Split Type:

Write-up: Patient c/o fatigue and cough on 1.4.2021 and was encouraged to be tested for COVID. We were notified that the patient was hospitalized on 1.7.2021 with COVID symptoms and positive test results. She is currently on a ventilator and dialysis.

VAERS ID: 928662 (history)
Form: Version 2.0
Age: 50.0
Sex: Female
Location: New York
Vaccinated: 2020-12-23

Onset: 2021-01-01
Days after vaccination: 9
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram abnormal, Arteriospasm coronary, Chest pain, Fibrin D dimer increased, Troponin I increased, Vasoconstriction
SMQs:, Haemorrhage laboratory terms (broad), Myocardial infarction (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Other ischaemic heart disease (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Synthroid Imitrex
Current Illness: None
Preexisting Conditions: Graves Disease Gerd
Allergies: Penicillin Sulfa drugs
Diagnostic Lab Data: January 3rd, Ddimer 763. Troponin .05, Troponin .06, January 5th Troponin .07, Troponin .03
CDC Split Type:

Write-up: On December 25th I had mild chest pain and then on January 1st, 2021 I had severe chest pain that persisted and on January 3rd I was admitted into the hospital. My Ddimer was elevated and my Troponin levels were elevated. An angiogram was performed and Dr. injected nitro into my arteries because they were constricted from Coronary Spasms.

VAERS ID: 928754 (history)
Form: Version 2.0
Age: 56.0
Sex: Female
Location: California
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00

Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1686 / 1	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Hypotension, Throat tightness
SMQs:, Anaphylactic reaction (narrow), Angioedema (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Vestibular disorders (broad), Hypersensitivity (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Sxs started 3-5 minutes post vax. Dizziness, hypotension, throat fullness, CODE called, given IM Epi at vax site. Taken to ED from vax site. Started on epi drip. Admitted to SHC.

VAERS ID: 928757 (history)
Form: Version 2.0
Age: 34.0
Sex: Male
Location: Illinois
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / -

Administered by: Public **Purchased by:** ?
Symptoms: Dizziness, Nausea, Syncope
SMQs:, Torsade de pointes/QT prolongation (broad), Acute pancreatitis (broad),

Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Na
Current Illness: Na
Preexisting Conditions: Na
Allergies: Na
Diagnostic Lab Data:
CDC Split Type:

Write-up: Nausea/ dizzy, Syncope 12 hours later.

VAERS ID: 928772 (history)
Form: Version 2.0
Age: 52.0
Sex: Female
Location: New York
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Activated partial thromboplastin time, Angiogram cerebral, Arteriogram carotid, Blood magnesium, Blood pressure increased, Chest X-ray, Culture urine, Differential white blood cell count, Dizziness, Electrocardiogram, Feeling hot, Full blood count, Glycosylated haemoglobin, International normalised ratio, Lipase, Lipids, Magnetic resonance imaging brain, Magnetic resonance imaging spinal, Metabolic function test, Palpitations, Presyncope, Prohormone brain natriuretic peptide, Prothrombin time, SARS-CoV-2 antibody test, SARS-CoV-2 test, Scan with contrast, Swallow study, Troponin, Urine analysis
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Hypertension (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode

(broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: GERD Migraine
Allergies: latex shellfish
Diagnostic Lab Data: EKG 12/30 & 12/31 MRI of head & cervical spine 12/30/2020 Chest x-ray 12/30 CT Angio of head & neck with contrast 12/30 12/30/2020 Urinalysis & urine culture A1C Troponin lipid profile Serum Pro-BNP PT/INR, PTT Magnesium lipase COVID-19 PCR COVID-19 Antibody CMP & BMP CBC with diff 12/31 Discharge screening
CDC Split Type:

Write-up: Patient felt warm with palpitations 5 minutes after vaccine administered. was monitored for 30 mins & then returned to work. on 12/30/2020 patient was at work in Presurgical testing dept & experienced near syncope, dizziness & elevated BP. reported to ED & was admitted to telemetry unit.

VAERS ID: 928777 (history)
Form: Version 2.0
Age: 49.0
Sex: Female
Location: Kentucky
Vaccinated: 2021-01-06
Onset: 2021-01-08
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Anaphylactic reaction, Injection site erythema, Pyrexia, Rash, Supraventricular tachycardia
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Supraventricular tachyarrhythmias (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Extravasation events (injections, infusions and implants) (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Dr. called this morning and reported that an employee that works in billing had her vaccine on Wednesday and developed and anaphylactic reaction to Moderna. This was 24 hours later with rash, SVT heart rate above 140, low grade fever, redness at site. Admitted and treated with steroids and Benadryl.

VAERS ID: 928792 (history)
Form: Version 2.0
Age: 56.0
Sex: Female
Location: Michigan
Vaccinated: 2020-12-18
Onset: 2020-12-20
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chills, Dyspnoea, Electrocardiogram ST segment elevation, Pain, Palpitations
SMQs:, Anaphylactic reaction (broad), Myocardial infarction (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Came to ER on 12/20/20 with chills, heart palpitations, body aches and increased SOB. Had ST elevation on EKG in ER, taken to Cath Lab- no intervention done. D/C home 12/22/20. Previous Hx of COVID per patient

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VAERS ID: 928871 (history)
Form: Version 2.0
Age: 56.0
Sex: Female
Location: California
Vaccinated: 2020-12-26
Onset: 2020-12-27
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Cough, Dyspnoea, Head discomfort, Oxygen saturation decreased, Paranasal sinus discomfort, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Respiratory failure (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Combipatch 0.05 mg/0.14mg/24 hrs transdermal film, 1 patch topical 2x per week olmesartan 20 mg daily
Current Illness:

Preexisting Conditions: hypertension
Allergies: penicillin--rash sulfa--rash
Diagnostic Lab Data: COVID-19 (SARS-CoV-2) Molecular Amp positive 12/29/20
CDC Split Type:

Write-up: Patient received first dose of Pfizer COVID-19 vaccine on December 26. On the next day, December 27, patient started having pressure in her head and sinuses, weakness. Then she developed nonproductive cough and progressive shortness of breath. She was seen at urgent care and tested positive for COVID-19 on December 29. She had low oxygen saturation on home oximeter and severe shortness of breath. Patient''s husband is also ill with COVID-19 at home. Patient was sent to the ED and admitted to the hospital.

VAERS ID: 929074 (history)
Form: Version 2.0
Age: 31.0
Sex: Female
Location: California
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J202A / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dysphagia, Lip swelling, Oropharyngeal pain, Palpitations, Speech disorder, Tachycardia
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ascorbic acid Vitamin D Vitamin E zinc sulfate
Current Illness: none
Preexisting Conditions: none

Allergies: phenazopyridine--hallucinations
Diagnostic Lab Data: Tachycardic with HR 120s, all other vital signs stable.
CDC Split Type:

Write-up: About 15 minutes after receiving the vaccine she felt palpitations. She was monitored for another 15 minutes and while she was walking to her car se started noticing sore throat associated with inability to talk, unable to swallow secretions, and swelling the lips. Patient presented to the emergency room where she received EpiPen dose. Received diphenhydramine, famotidine, and prednisone. Lip swelling and sore throat began improving in ED.

VAERS ID: 929184 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-29
Onset: 2021-01-01
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / UNK	RA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Dysphagia, Dyspnoea, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Anxiety (Diagnosed 22 years ago)
Preexisting Conditions:
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021006254

Write-up: difficulty breathing/swallowing; difficulty breathing/swallowing; Hives/hives all over including in her mouth; This is a spontaneous report from a contactable consumer (spouse). A 46-year-old female patient (wife) received BNT162B2 (Pfizer-Biontech Covid-19 Vaccine, lot number: EH9899) via unspecified route of administration at arm right on 29Dec2020 at single dose for Covid-19 vaccine. Medical history included ongoing anxiety diagnosed 22 years ago. Concomitant medications were not none. Patient received the vaccine and was one big hive (01Jan2021). She went to the ER on sat and will have to go back. Caller is asking where she should go. patient went through the VAERS report 3 times. She did not have any sides effects. patient is a frontline worker. On 29Dec2020 she got the Covid vaccine. On Sat 02Jan2021 she went the ER with hives all over including in her mouth, stated she had difficulty breathing/swallowing, was given medication to take home and discharged. She was readmitted yesterday 05Jan2020 with worsening symptoms and needed to be given a prednisone nebulizer. they had two ER visits. No Investigation Assessment. patient was not recovered from the event hives/hives all over including in her mouth, the final outcome of other events was unknown.

VAERS ID: 929391 (history)
Form: Version 2.0
Age: 37.0
Sex: Female
Location: New York
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Acute respiratory failure, Anaphylactic reaction, Dysphagia, Endotracheal intubation, Heart rate increased, Swelling of eyelid, Vomiting
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Hypoglycaemic and neurogenic shock conditions (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Periorbital and eyelid disorders (narrow), Hypersensitivity (narrow), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No

Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Escitalopram 10mg 1 po Qday Gabapentin 300mg po three times a day
Current Illness:
Preexisting Conditions: Bell"s palsy Migraine Seizures Systemic lupus erythematosus
Allergies: Dilantin Ibuprofen Latex Morphine Penicillins Tegretol Valium
Diagnostic Lab Data:
CDC Split Type:

Write-up: 1/6/21 Pt received vaccine and complained of difficulty swallowing and rapid heart rate. Pt received methylprednisolone 125mg IVP, diphenhydramine 25mg IVP, & famotidine 20mg IVP. Pt reported improvement and was discharged. Sent home on diphenhydramine and oral prednisone. 1/7/21 Pt unable to swallow her own secretions and experienced eyelid swelling. Pt vomitted. Pt received epinephrine and Benadryl X 1 dose each. Pt then transported to hospital via ambulance. Reason for admission - acute respiratory failure secondary to anaphylactic reaction. Decision was made to emergently intubate the patient for airway protection despite aggressive intervention. Pt successfully extubated 1/8/21. Plan to discharge home and start Medrol Dose Pack 1/9/21.

VAERS ID: 929610 (history)
Form: Version 2.0
Age: 83.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Align, Amlodipine, Arimidex, Atorvastatin, Calcium, Colace, Gabapentin, Losarten, Mucinex, Namenda, MVI, Pataday drops, Prilosec, Remeron, Vitamin d3
Current Illness: Pneumonia (resolved 12-17-20
Preexisting Conditions: COPD, hypertension, PVD, neuropathy, RA, GERD, OP, breast cancer
Allergies: PCN, Shellfish
Diagnostic Lab Data:
CDC Split Type:

Write-up: C/o shortness of breath routine oxygen increased cannula changed to mask oxygen sats at 88%

VAERS ID: 929683 (history)
Form: Version 2.0
Age: 93.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Dyspnoea, Muscular weakness
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Peripheral neuropathy (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: Levothyroxine, Lisinopril, Metoprolol, Prozac, Trazadone
Current Illness: Pneumonia, UTI with ESBL
Preexisting Conditions: A-Fib, Cardiomegaly, HTN, CHF, HLD, Hypothyroidism
Allergies: NKA
Diagnostic Lab Data: none transferred to hospital
CDC Split Type:

Write-up: Labored breathing with oxygen running at 4l/min, muscle weakness

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VAERS ID: 929689 (history)
Form: Version 2.0
Age: 61.0
Sex: Male
Location: California
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 2	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Bacteraemia, Blood culture, Blood lactic acid, Culture urine, Full blood count, Laboratory test, Liver function test, Pyrexia, Respiratory viral panel, SARS-CoV-2 test negative, Urine analysis normal, Urosepsis
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Sepsis (narrow), Opportunistic infections (broad), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: HCTZ, losartan, carvedilol, aspirin
Current Illness: none
Preexisting Conditions: h/o multiple strokes, vascular dementia (bedbound, unable to speak) trach for h/o resp failure, G-tube/tubefeed dep
Allergies: amlodipine (gum swelling)
Diagnostic Lab Data: Cultures (blood, urine), COVID PCR & Biofire resp panel on nasopharyngeal swab (negative), CBC, Chem10, lactate, LFTs

CDC Split Type:

Write-up: Fever to 103.7F, respiratory rate 36. Was transferred from facility to hospital. Since then has been found to have gram-negative rod bacteremia, although urinalysis was negative, urine culture pending. Patient has since defervesced after receiving 1 dose of cefepime. Overall the most likely cause of fever seems to be urosepsis w/ bacteremia, pending confirmation with urine & blood cultures.

VAERS ID: 930005 (history)
Form: Version 2.0
Age: 26.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-22
Onset: 2020-12-22
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Chest X-ray, Computerised tomogram head, Headache, Hypoaesthesia, Lumbar puncture, Magnetic resonance imaging spinal, Metabolic function test, Paraesthesia, Swollen tongue, Syncope, Urine analysis
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (broad), Angioedema (narrow), Peripheral neuropathy (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 6 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none
Preexisting Conditions: cholecystectomy one month prior to vaccination with chronic diarrhea since
Allergies: cephalosporins, tramadol
Diagnostic Lab Data: head CT, cxr, spinal mri, spinal tap, cmp/vitals/UA

CDC Split Type:

Write-up: initial; swelling of tongue, tingling and numbness in legs, syncope. later; HA

VAERS ID: 930010 (history)
Form: Version 2.0
Age: 87.0
Sex: Male
Location: Minnesota
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Asthenia, Blood pressure increased, COVID-19 pneumonia, Confusional state, Dyspnoea, Fatigue, Feeling abnormal, Hypoxia, Influenza A virus test negative, Influenza B virus test, Influenza virus test negative, Intensive care, Pyrexia, Respiratory syncytial virus test negative, SARS-CoV-2 test positive, Tremor
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Parkinson-like events (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypertension (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Citrucel, acetaminophen 8 hour, systane ultra, gabapentin, pantoprazole sodium, vitamin c, vitamin d3, aspirin, atorvastatin calcium, cymbalta, norco, lantus solostar, loperamide, melatonin, toprol xl, flomax, preservision areds, vitamin b-
Current Illness:
Preexisting Conditions: Peripheral vascular disease, chronic systolic heart failure, type 2 diabetes mellitus, essential hypertension, hyperlipidemia, chronic kidney disease stage 3 unspecified, gastro-esophageal reflux disease without esophagitis, hypothyroidism, acquired absence of left and right legs below the knee
Allergies: Colestipol, Amitriptyline, Colestid, Niacin, Ondansetron

Diagnostic Lab Data: COVID-19 PCR test 1/5/21 negative COVID-19 PCR test 1/6/21 positive Influenza A, B; RSV negative 1/6/21
CDC Split Type:

Write-up: Resident displayed with confusion/shaking at 1400, condition worsens at time went. Resident unable to state where he is, knows his name. can tell you he does not feel right. Temp 97.3, p 88, O2 91%, Bp 214/116 Transferred to ED with fever, temp of 103, and shortness of breath, admitted to ICU Positive COVID-19 test at hospital. Diagnoses include acute COVID-19 pneumonia and hypoxia. PO had confusion, fatigue, weakness, hypoxia, increased BP

VAERS ID: 930109 (history)
Form: Version 2.0
Age: 93.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3246 / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Dizziness, Ear pain
SMQs:, Anticholinergic syndrome (broad), Vestibular disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Insulin, Jardiance, Januvia, Levemir, Novalog, Lipitor
Current Illness: kidney failure
Preexisting Conditions: Diabetes
Allergies: nokidney failure
Diagnostic Lab Data: no
CDC Split Type:

Write-up: She began with an earache and dizziness. Pain got so severe that she could no longer take it. Went to the doctor which she was put on pain medications. Went to ER on 1/6

and on 1/7 went to her PCP. Still in severe pain.

VAERS ID: 930142 (history)
Form: Version 2.0
Age: 68.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-05
Onset: 2021-01-08
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / IM

Administered by: Other **Purchased by:** ?
Symptoms: Chest pain, Stress cardiomyopathy, Troponin increased
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (narrow), Other ischaemic heart disease (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Nortriptline Multivitamin
Current Illness:
Preexisting Conditions:
Allergies: No Known Allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient experiencing Chest pain and elevated troponin. Patient taken to the cath lab and treated for suspected stress induced cardiomyopathy.

VAERS ID: 930153 (history)
Form: Version 2.0
Age: 41.0
Sex: Male

Location: Michigan
Vaccinated: 2021-01-05
Onset: 2021-01-07
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Condition aggravated, Immune thrombocytopenia, Platelet count decreased
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Atorvastain 10mg Levothyroxine 112mcg Prilosec 20mg Insulin
Current Illness: NA
Preexisting Conditions: Insulin dependent Diabetes HX of ITP 2014, in remession
Allergies: PCN
Diagnostic Lab Data: Plt 2
CDC Split Type:

Write-up: ITP Plt 2

VAERS ID: 930235 (history)
Form: Version 2.0
Age: 44.0
Sex: Male
Location: Colorado
Vaccinated: 2020-12-30
Onset: 2021-01-02
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Abdominal pain, Asthenia, Back pain, Computerised tomogram spine, Muscular weakness, Paraesthesia, Reflexes abnormal
SMQs:, Rhabdomyolysis/myopathy (broad), Acute pancreatitis (broad), Peripheral neuropathy (broad), Retroperitoneal fibrosis (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Multivitamin daily Ambien 5mg daily/nighttime
Current Illness:
Preexisting Conditions: History of Bells Palsy
Allergies:
Diagnostic Lab Data: CT L-Spine *NSMC 01/02
CDC Split Type:

Write-up: Hospital on 1/2 - then again on 1/5, transferred and admitted to hospital, discharged 1/6 Abnormal reflex/weakness back pain paresthesia and weakness of legs abdominal pain evaluation for possible GBS post covid 19 vaccine

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VAERS ID: 930297 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Rhode Island
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood creatine phosphokinase normal, Blood lactic acid, Computerised tomogram head normal, Dysphonia, Insomnia, Muscle rigidity, Nausea, Oropharyngeal pain, Retching, Speech disorder, Tremor, Vomiting

SMQs:, Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Dementia (broad), Parkinson-like events (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Adderall
Current Illness:
Preexisting Conditions: ADHD
Allergies: Adhesive tape (NOT allergic to latex)
Diagnostic Lab Data: CT scan, lactate, CPK normal
CDC Split Type:

Write-up: 29-year-old previously healthy female presenting today with difficulty sleeping, sore throat, and nausea after receiving the second Pfizer COVID-19 vaccine around 10 AM. Patient says that after her first dose of the vaccine she had mild sore throat and hoarse voice that resolved spontaneously. She had her vaccine around 10, several hours later before coming to the emergency department between 2--230 she had the sudden onset of difficulty speaking with associated sore throat and nausea. She has had dry heaving but no large amounts of vomiting. She has not had stridor, wheezing, shortness of breath, syncope, or the development of a rash or hives. She has not had a reaction to her prior vaccines she does not have any other allergies in general. She has otherwise been well recently without infectious symptoms including fevers, chills, cough, and has not been exposed to Covid to her knowledge. Medications administered in ED included diphenhydramine 50 mg IV once, dexamethasone 10 mg IV once, famotidine 40 mg IV once, ondansetron 4 mg IV once. Had brief episode of shaking and R arm rigidity in the ED. Patient denies LOC during the episode, did not have a post ictal state, no tongue biting or episodes of incontinence. Given single event without LOC, less concern about episode being a seizure. Patient has had no further episodes of shaking since admission. CT brain was wnl, no FNDs noted on exam.

VAERS ID:	930348 (history)
Form:	Version 2.0
Age:	33.0
Sex:	Female
Location:	New Jersey
Vaccinated:	2020-12-28
Onset:	2021-01-01
Days after vaccination:	4

Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / SYR

Administered by: Senior Living **Purchased by:** ?
Symptoms: Maternal exposure during breast feeding
SMQs:, Neonatal exposures via breast milk (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Prenatal, sunflower lecithin
Current Illness: No
Preexisting Conditions: Pcos
Allergies: Tree nuts and shellfish
Diagnostic Lab Data: Video eeg starting on 1/6-1/8. Lab work on my daughter. All clear. Episodes stopped after 1/5.
CDC Split Type:

Write-up: I am breastfeeding. My daughter had seizure like episodes starting on Saturday 1/2, Sunday 1/3, Monday, 1/4 and 2 times on Tuesday 1/5.

VAERS ID: 930476 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3246 / 2	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Differential white blood cell count, Full blood count, Glomerular filtration rate, Hyperhidrosis, Hypersensitivity, Loss of consciousness, Metabolic function test, Pharyngeal

erythema, Pharyngeal swelling, Platelet count, Rash, Swelling face, Swollen tongue
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (narrow), Angioedema (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Lysine Zoloft Adzenys Evekeo Vitamin d Docusate sodium
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: CBC with platelet and differential x 2 Basic Metabolic Panel Estimated GFR
CDC Split Type:

Write-up: Acute allergic reaction Tongue swelling Facial swelling Throat swelling Rash on throat and chest Redness in throat Diaphoresis Momentary loss of consciousness

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VAERS ID: 930484 (history)
Form: Version 2.0
Age: 35.0
Sex: Female
Location: Wyoming
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1686 / 1	AR / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Mental status changes
SMQs:, Dementia (broad), Noninfectious encephalitis (broad), Noninfectious

encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness: Unknown
Preexisting Conditions: Unknown
Allergies: Unknown
Diagnostic Lab Data: Unknown
CDC Split Type:

Write-up: Altered Mental Status began the middle of the night of 01/06/21 and 01/07/21 with worsening overall status- definitive symptoms unknown to this reporter, this person was admitted to the local hospital at approximately 1800 01/07/2021. This reporter was not told the admitting diagnosis or any defining symptoms, only that the person was admitted.

VAERS ID: 930508 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: California
Vaccinated: 2021-01-08
Onset: 2021-01-08
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	9899 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dyspnoea, Hypotension, Injection site pruritus, Rash, Tachycardia, Throat irritation, Urticaria
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? Yes

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Phentermine (ADIPEX-P) 37.5 mg Oral Cap Hydrocortisone 2.5 % Top Crea cloNIDine HCL (CATAPRES) 0.2 mg Oral Tab tiZANidine (ZANAFLEX) 4 mg Oral Tab traZODone (DESYREL) 50 mg Oral Tab Multivitamin (MULTIPLE VITAMIN) Oral Cap
Current Illness: unknown
Preexisting Conditions:
Allergies: Acetaminophen-codeine Cymbalta [Duloxetine Hcl]
Diagnostic Lab Data:
CDC Split Type:

Write-up: Initial itching at injection site, observed and returned to work. Came back ~30-40 minutes later with itchiness in throat and hives to arm. Given Benadryl PO and observed for extended period of time. Symptoms not resolving. Patient transferred to Emergency Department for further care. At that point observed to have full body rash, SOB. Given Epi while in ED. Developed tachycardia, hypotension. Treatment continued.

VAERS ID: 930518 (history)
Form: Version 2.0
Age: 33.0
Sex: Male
Location: Maryland
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Blood potassium decreased, Blood pressure abnormal, Dehydration, Echocardiogram normal, Electrocardiogram normal, Fatigue, Feeling abnormal, Flushing, Headache, Heart rate increased, Peripheral swelling, Pulse abnormal
SMQs:, Cardiac failure (broad), Anaphylactic reaction (broad), Angioedema (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypertension (broad), Hypersensitivity (broad), Dehydration (narrow), Hypokalaemia (narrow)

Life Threatening? No
Birth Defect? No

Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Dupuytrens, plantar fibromatosis, peyronies
Allergies: None
Diagnostic Lab Data: Low potassium, cardio gram was normal, ekg was normal just very rapid beat, I was told most of my lands were great, had signs of mild dehydration.
CDC Split Type:

Write-up: After about 15 min: hr went to over 170bpm , flushed sensation, brain fog, was driving at the time and my brain wouldn?t figure out how to call 911 , eventually I figured it out but it took extreme mental effort. Paramedics came and my pulse was From 100 - 170 changing rapidly, bp 150-175/x changing rapidly, symptoms would come and go about 8 times on way to hospital. In ER same thing then about 2 hrs go by and then all started again. Pulse remained at 90- 100bpm at least until 2am. (My resting hr is between 40 and 50, my normal bp is usually sub 120/80) I later realized my hands were swollen, mostly on my left side. Lingering symptoms include brain fog, tiredness, and headache. Maybe an exasperating issue was I had low potassium levels likely making symptoms worse - I worked out heavily early in the morning, probably had low intake of potassium that day and day before, and maybe slightly dehydrated from my workout - I do Functional lifting, run, and other types of exercise regularly.

VAERS ID: 930611 (history)
Form: Version 2.0
Age: 71.0
Sex: Male
Location: Massachusetts
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Bilevel positive airway pressure, Cardiac failure congestive, Condition aggravated, Intensive care, Respiratory failure
SMQs:, Cardiac failure (narrow), Anaphylactic reaction (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (narrow), Guillain-Barre

syndrome (broad), Cardiomyopathy (broad), Hypersensitivity (broad), Respiratory failure (narrow), Hypokalaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Albuterol, metoprolol, fish oil, apixiban, insulin
Current Illness: Advanced diastolic heart failure and CKD. Returned from hospitalization one week prior for CHF exacerbation treated with BIPAP, IV diuresis
Preexisting Conditions: oxygen dependent COPD, type II DM, diastolic CHF, stage IV renal failure, personality disorder
Allergies: statin, PCN
Diagnostic Lab Data: unknonw
CDC Split Type:

Write-up: Developed hypercapnic respiratory failure, CHF exacerbation - readmitted to Hospital. In ICU with BIPAP

VAERS ID: 930669 (history)
Form: Version 2.0
Age: 56.0
Sex: Male
Location: Arkansas
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	PAA156051 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dehydration, Feeling abnormal, Malaise, SARS-CoV-2 test positive
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Dementia (broad), Infective pneumonia (broad), Dehydration (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none reported
Current Illness: none reported
Preexisting Conditions: none reported
Allergies: none reported
Diagnostic Lab Data: COVID-19 test positive
CDC Split Type:

Write-up: patient begin to feel bad that night was admitted into hospital sometime in the next couple of days for dehydration, patient discharged home and then readmitted to hospital for positive covid testing after feeling very ill.

VAERS ID: 930889 (history)
Form: Version 2.0
Age: 48.0
Sex: Male
Location: Minnesota
Vaccinated: 2020-12-22
Onset: 2020-12-27
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angioplasty, Coronary artery occlusion, Electrocardiogram ST segment elevation, Myocardial infarction, Stent placement
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Propranolol, Pepcid,cetrizine, losartan
Current Illness: None

Preexisting Conditions: HTN, allergies
Allergies: Nkda
Diagnostic Lab Data: I had a myocardial infarction on December 27, 2020. I had received my first vaccination for COVID-19 on December 22, 2020. I had a ST segment elevation MI requiring emergent angio plasty and stenting with a severe ?99%? proximal LAD lesion.
CDC Split Type:

Write-up: I had a myocardial infarction on December 27, 2020. I had received my first vaccination for COVID-19 on December 22, 2020. Not sure if these are related but I felt I should report it.

VAERS ID: 930894 (history)
Form: Version 2.0
Age: 75.0
Sex: Male
Location: Guam
Vaccinated: 2020-12-29
Onset: 2021-01-02
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	AR / -

Administered by: Private **Purchased by:** ?
Symptoms: Blood pressure increased, Brain herniation, Computerised tomogram abnormal, Computerised tomogram head, Haemoglobin increased, Haemorrhage intracranial, Headache, Hypertension, Hypertensive emergency, Hypotonia, Intensive care, International normalised ratio normal, Intracerebral haematoma evacuation, Platelet count normal, Pyrexia, Shift to the left, White blood cell count normal
SMQs:, Peripheral neuropathy (broad), Haemorrhage terms (excl laboratory terms) (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Haemorrhagic central nervous system vascular conditions (narrow), Guillain-Barre syndrome (broad), Hypertension (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Not on medications

Current Illness: Hypertension
Preexisting Conditions: Hypertension
Allergies: Possibly penicillin
Diagnostic Lab Data: Initial BP 178/73 crept up to $g200 despite medications, initially controlled then difficult to control SBP 160-211 Noted new flacidity 10am 1/2/2021 Stat CT 1/2/2021 head Acute intracranial hemorrhage 6.5x6x 5 cm with beginning of herniation , left shift Labs on 1/1/2021 WBC 8.8, Hgb 16, platelets 160 INR 1/2 =1.0
CDC Split Type:

Write-up: Low grade Fever, headache needing admission Intracranial hemorrhage with hypertension Medical management for hypertensive emergency Received surgical evacuation admitted in Intensive care,

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VAERS ID: 950259 (history)
Form: Version 2.0
Age: 58.0
Sex: Female
Location: Unknown
Vaccinated: 2020-12-23
Onset: 2020-12-24
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Abdominal pain, Appendicectomy, Appendicitis, Nausea
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Appendectomy Narrative: Developed abdominal pain with nausea on 12/24/2020,

went to ER and noted to have appendicitis, resulting in an appendectomy

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VAERS ID: 928990 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-17
Onset: 2020-12-19
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Ischaemic stroke
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Aortic aneurysm repair; Gout; Hypertension; Middle cerebral artery infarct (Left MCA infarct); Middle cerebral artery stroke; Peripheral vascular disease
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021004018

Write-up: Potential ischaemic stroke; This is a spontaneous report from a contactable physician downloaded from the Agency Regulatory Authority-WEB. Regulatory authority GB-MHRA-EYC 00235997, Safety Report Unique Identifier GB-MHRA-ADR 24545778. An 81-year-old male patient received bnt162b2 (COMIRNATY, lot number: unknown, strength: 30mcg/0.3ml) intramuscular on 17Dec2020 at 0.3 mL, single for COVID-19 vaccination (covid-19 immunisation). Medical history included hypertension from an unknown date and unknown if ongoing, aortic aneurysm repair in 2008 (not ongoing), peripheral vascular disorder from an unknown date and unknown if ongoing, peripheral vascular disease from an unknown date and unknown if ongoing, middle cerebral artery infarct from 2019 to an unknown date and

unknown if ongoing (Left MCA infarct), gout from an unknown date and unknown if ongoing, middle cerebral artery stroke from an unknown date and unknown if ongoing. Concomitant medication included paracetamol, lansoprazole, clopidogrel, perindopril, allopurinol, atorvastatin, bisoprolol. The patient experienced potential ischaemic stroke (hospitalization, disability) on 19Dec2020. It was reported presented with ischaemic stroke, National Institutes of Health Stroke Scale (NIHSS 30) two days following vaccination. Risk factors for patient include previous stroke, peripheral vascular disease (PVD). Thrombosed for left middle cerebral artery (MCA) stroke 19Dec2020. Patient was not recovered from the event. No follow-up attempts possible. No further information expected. Information on lot and batch numbers cannot be obtained.

VAERS ID: 928992 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-18
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Atrial fibrillation, Condition aggravated, Death, Malaise
SMQs:, Supraventricular tachyarrhythmias (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2020-12-20
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ;
Current Illness: Atrial fibrillation; Diabetes; Frailty; Hypothyroidism; Osteoporosis
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021003922

Write-up: Atrial fibrillation; This is a spontaneous report from a contactable physician downloaded from the Regulatory Authority. The regulatory authority report number is GB-MHRA-EYC 00236011. An 87-year-old female patient received BNT162B2 (PFIZER-

BIONTECH COVID-19 mRNA VACCINE; Lot number EJ0553), intramuscular on 18Dec2020 at 0.3 mL, single for covid-19 immunization. Medical history included ongoing hypothyroidism, ongoing diabetes, ongoing atrial fibrillation, ongoing frailty and, ongoing osteoporosis, all from unknown dates. Concomitant medication included prednisolone (MANUFACTURER UNKNOWN), levothyroxine (MANUFACTURER UNKNOWN), salbutamol (MANUFACTURER UNKNOWN), omeprazole (MANUFACTURER UNKNOWN), doxycycline (MANUFACTURER UNKNOWN). The patient experienced atrial fibrillation on an unspecified date, which was serious as it was medically significant, involved hospitalization and lead to death. Clinical course was as follows: the patient was vaccinated. Consent was obtained and a pre immunization checklist was completed. She was observed following the administration of the vaccine, and no adverse effects were noted. She returned home. She became unwell and was admitted to hospital approximately 24 hours later. The patient was admitted to the hospital 24 hours following the vaccination, and subsequently died later, while in the hospital. The full clinical details were unknown, but the diagnosis from Accident & Emergency was atrial fibrillation. It is not clear if this had any relation to the vaccine that was administered, but could not be excluded, per the reporter. The patient died on 20Dec2020. It was not reported if an autopsy was performed. No follow-up activities are possible. No further information is expected.; Reported Cause(s) of Death: Atrial fibrillation

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VAERS ID: 929011 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Dizziness, Heart rate, SARS-CoV-2 test, Syncope

SMQs:, Torsade de pointes/QT prolongation (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days

Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201221; Test Name: Blood pressure; Result Unstructured Data: Test Result:171/71; Test Date: 20201221; Test Name: Pulse rate; Result Unstructured Data: Test Result:65; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2021003966

Write-up: Dizzy; Syncope vasovagal; This is a spontaneous report from a contactable physician downloaded from the Agency Regulatory Authority-WEB. The regulatory authority report number GB-MHRA-WEBCOVID-20201221152032. A male patient of an unspecified age received BNT162B2 (COMIRNATY- PFIZER-BIONTECH COVID-19 mRNA VACCINE; Lot number EJ0553), via an unspecified route of administration on 21Dec2020 as single dose for COVID -19 immunization. The patient"s medical history was not reported. Concomitant medication included bisoprolol (MANUFACTURER UNKNOWN) for atrial fibrillation. The patient previously took naproxen and experienced unspecified adverse drug reaction, and benzocaine, salicylamide (INTRALGIN) and experienced unspecified adverse drug reaction. The patient was dizzy and experienced syncope vasovagal on 21Dec2020, which were both serious per hospitalization, was medically significant and life-threatening. The patient underwent lab tests and procedures which included blood pressure: 171/71 on 21Dec2020, pulse rate: 65 on 21Dec2020, COVID-19 virus test: negative on an unspecified date. Patient was not enrolled in a clinical trial. Patient has not tested positive for COVID-19 since the vaccine. Patient was driving home following his vaccination at the center. The patient had waited the required 15 minutes and felt well, so he left. Whilst driving home, the patient described feeling dizzy and lightheaded and then there was a loud bang and he has crashed his car. On attendance in accident and emergency, his blood pressure was 171/71 pulse was 65, he was not confused or disorientated, and no reduction of consciousness was noted. No issues with his airway or breathing, no skin or mucosal changes, no signs of anaphylaxis were noted. Following discussion with the doctor in accident and emergency, it was felt the likely diagnosis was a vasovagal episode. Patient was observed for a period of time following which he was discharged. No follow-up attempts are possible. No further information is expected.

VAERS ID: 929025 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?

Symptoms: Consciousness fluctuating, Dizziness, Somnolence, Tremor
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Parkinson-like events (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Vestibular disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021005230

Write-up: Received covid vaccine and once leaving observation room experienced light headed, shaking and drowsiness / varying levels of consciousness.; shaking; drowsiness; varying levels of consciousness; This is a spontaneous report from a contactable pharmacist received via a sales representative. A female patient of an unspecified age received single dose of BNT162B2 (Solution for injection, batch/lot and exp date not reported), via unspecified route of administration on an unspecified date for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. The pharmacist reported that the patient received the Covid vaccine on an unspecified date and once leaving the observation room experienced lightheadedness, shaking, drowsiness, and varying levels of consciousness. The events were considered as serious due to hospitalization. The pharmacist reported that the patient was sent home and was well. The patient recovered from the events on an unspecified date. No follow-up attempts are possible; information about batch number cannot be obtained.; Sender"s Comments: A possible contributory effect of suspect BNT162B2 on reported events cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID:	929028 (history)
Form:	Version 2.0
Age:	
Sex:	Male
Location:	Foreign
Vaccinated:	2020-12-30
Onset:	2021-01-01

Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Death, Endotracheal intubation, Pleural effusion, Pyrexia, Respiratory distress, Sepsis, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (broad), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Acute central respiratory depression (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Haemodynamic oedema, effusions and fluid overload (narrow), Hypersensitivity (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Sepsis (narrow), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-04
Days after onset: 3
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Bladder cancer; Blood pressure abnormal; Diabetes
Allergies:
Diagnostic Lab Data:
CDC Split Type: ILPFIZER INC2021009752

Write-up: SEPSIS; respiratory distress; PLEURAL EFFUSION; This is a spontaneous report received from other healthcare professional via the Division of epidemiology of the Ministry of Health. The other healthcare professional reported similar events for three patients. This is the third of three reports. A 91-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 30Dec2020 at single dose for covid-19 immunisation. Medical history included known background of blood pressure disease, diabetes, malignant bladder from an unknown date and unknown if ongoing. The patient''s concomitant medications were not reported. Patient was received at the emergency room 3 days after receiving the corona vaccine in Jan2021, with fever, vomiting more than 40 times, in respiratory distress, was hospitalized in internal medicine department with sepsis diagnosis due to respiratory distress and pleural effusion, intubated, his condition was serious, patient passed away on 04Jan2021. Cause of death was reported as sepsis, respiratory distress and pleural effusion. It was not reported if an autopsy was performed. Follow-up attempts are completed. No further information is expected. Information about batch/lot

number cannot be obtained.; Sender"s Comments: Based on the information currently provided, the fatal events sepsis, respiratory distress and pleural effusion are more likely attributed to intercurrent infectious conditions associated with the advanced old patient underlying diseases . The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.,Linked Report(s) : IL-PFIZER INC-2020519349 same reporter, product, similar event, different patient;IL-PFIZER INC-2021009751 same reporter, product, similar event, different patient; Reported Cause(s) of Death: SEPSIS; respiratory distress; PLEURAL EFFUSION

VAERS ID: 929034 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6796 / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Anxiety, Asthenia, Blood pressure increased, Blood pressure measurement, Body temperature, Body temperature increased, Cold sweat, Dry mouth, Dysgeusia, Peripheral coldness, Syncope
SMQs:, Torsade de pointes/QT prolongation (broad), Neuroleptic malignant syndrome (broad), Taste and smell disorders (narrow), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Hypertension (narrow), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: BETAC; ;

Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20210105; Test Name: blood pressure; Result Unstructured Data: Test Result:170/100 mmHg; Comments: around 8 AM; Test Date: 20210105; Test Name: blood pressure; Result Unstructured Data: Test Result:160/100 mmHg; Comments: at 12 PM; Test Date: 20210104; Test Name: body temperature; Result Unstructured Data: Test Result:37.5 Centigrade
CDC Split Type: LVPFIZER INC2021006858

Write-up: blood pressure at this time was 170/100 mmHg; skin was cold; feeling near fainting; Cold sweat; anxiety; weakness; increased body temperature (37.5 C); dry mouth; bitter taste in mouth; This is a spontaneous report from a contactable physician. This is a report received from the Regulatory Authority. Regulatory authority report number LV-SAM-2021014370. A 46-years-old female patient received bnt162b2 (COMIRNATY), intramuscular on 04Jan2021 at 0.3 mL, single (Lot # EJ6796) for covid-19 immunisation. The patient medical history was not reported. Concomitant medication included betaxolol hydrochloride (BETAC), olmesartan, amlodipine. The patient experienced feeling near fainting, cold sweat, anxiety, weakness, increased body temperature (37.5 c), dry mouth, bitter taste in mouth, all on 04Jan2021; and blood pressure at this time was 170/100 mmhg on 05Jan2021. The patient was hospitalized for the events from 05Jan2021. Clinical course reported as: The patient who was vaccinated on 04Jan2021 and 20 minutes after receiving a dose of the bnt162b2, was driving a car and experienced a feeling of near fainting, accompanied by a cold sweat. After arriving home the patient still felt faint, had anxiety, weakness and increased body temperature (37.5 C), as well as dry mouth and bitter taste in mouth that decreased around 6 PM. On 05Jan2021 around 8 AM, the patient experienced a worsening of symptoms (ADR''s). Patient''s blood pressure at this time was 170/100 mmHg. At the time of the report 05Jan2021 at 12PM the patient''s blood pressure was 160/100 mmHg with sustained feeling of weakness, the patient''s skin was cold. The patient was hospitalized and her symptoms improved after receiving dexamethasone (dose unspecified). Out come of events was recovering.

VAERS ID: 930916 (history)
Form: Version 2.0
Age: 32.0
Sex: Female
Location: New Mexico
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Exposure during pregnancy, Premature baby, Premature delivery, Vaginal haemorrhage
SMQs:, Acute pancreatitis (broad), Haemorrhage terms (excl laboratory terms) (narrow), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic

procedures (narrow), Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow), Neonatal disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: prenatal vitamin calcium supplement 1200 mg vitamin d3 1000 iu Aspirin 81 mg
Current Illness: none
Preexisting Conditions: none
Allergies: NKA
Diagnostic Lab Data: see above
CDC Split Type:

Write-up: Patient is a 32 yo G2P1001 with EDD 5/2/2021 by 7w US. She had the first dose of the Pfizer Covid 19 vaccination on 12/17/2020 at the Health Clinic and the second dose on 1/7/2021 at 1115 am. She began having abdominal pain and vaginal bleeding at 315 sm on 1/8/2021 progressing to a previable (22w2d) preterm birth at 739pm on 1/8/2021.

VAERS ID: 931224 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-04
Onset: 2021-01-05
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1686 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Blood thyroid stimulating hormone normal, CSF cell count normal, CSF culture, CSF protein normal, Computerised tomogram head, Computerised tomogram head normal, Fatigue, Full blood count normal, Gait disturbance, Headache, Hyporeflexia, Lumbar puncture normal, Metabolic function test normal, Muscular weakness, Somnolence, Speech disorder, Spinal X-ray normal, Syncope, Vitamin B12 decreased, Vomiting
SMQs:, Torsade de pointes/QT prolongation (broad), Rhabdomyolysis/myopathy (broad), Acute pancreatitis (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Parkinson-

like events (broad), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Levothyroxine 200 mcg, Vyvanse 60 mg
Current Illness: NONE
Preexisting Conditions: Hypothyroidism, ADHD
Allergies: SHELLFISH, CECLOR
Diagnostic Lab Data: 1/6/21 Normal BMP, CBC, normal TSH, low B12 Lumbar Puncture: normal cell count and protein, culture pending Cervical Spine xray: Normal CT brain (noncontrast): normal
CDC Split Type:

Write-up: Pt experienced extreme fatigue and sleepiness the day following her second vaccination for Covid 19 and was found by her family after collapsing on 1/6/21 at 05:30. Upon arousal, she experienced headache, vomiting, weakness, difficulty speaking and difficulty walking with lower extremity weakness. She was taken to urgent care and subsequently admitted for evaluation at hospital and found to have a normal chemistry, blood count, normal lumbar puncture and normal imaging of her neck and brain. Discharge summary notes 3/5 strength and hyporeflexia throughout. Pt had televisit consult with psychiatry and neurology. She is subsequently to be discharged to a Facility without explanation for her sudden onset of progressive lower extremity and vocal weakness. She is noted to have a history of shellfish allergy. She experienced mild symptoms after the first vaccination, but no neurologic or vascular symptoms at that time.

VAERS ID: 931286 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Massachusetts
Vaccinated: 2020-01-04
Onset: 2021-01-05
Days after vaccination: 367
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Audiogram abnormal, Auditory disorder, Balance disorder, Blood test normal, Deafness, Deafness bilateral, Electrocardiogram normal, Fatigue, Magnetic resonance imaging brain normal, Myalgia, Pyrexia, Tinnitus, Vertigo
SMQs:, Rhabdomyolysis/myopathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Eosinophilic pneumonia (broad), Hearing impairment (narrow), Vestibular disorders (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Topiramate, Nortriptline, ibuprofen, Excedrin migraine, Zolmitriptan, Estrace cream
Current Illness: none
Preexisting Conditions: History of Graves disease, history of episodic vertigo with hearing loss thought to be due to inflammatory hair cell disease
Allergies: Sulfa, SSRI
Diagnostic Lab Data: brain MRI negative including auditory canals, blood work and EKG negative, audiogram showing profound hearing loss more at high frequencies and loss of discrimination in R ear.
CDC Split Type:

Write-up: I am a physician and I got dose 2 at 1:30pm on Jan 4. Next afternoon, Jan 5, I got severe myalgias, fever up to 100, severe fatigue, went home after work and slept til the next morning, went to work, took ibuprofen, and the myalgias improved and felt better. But around 3 pm, Jan 6, I got mild vertigo. By about 7pm Jan 6, I noticed my L ear didn"t hear well. I changed the battery in my hearing aide and cleaned it but It made no difference. I woke up on Jan 7 with severe vertigo and hearing loss. I did Epley"s maneuvers with no effect. I have had similar episodes. I went to work, but gave up when I could not hear patients talking to me. I went to the Emergency Dept and got admitted. I was too unsteady on my feet. Audiogram showed profound hearing loss both ears and almost complete loss of discrmination in R ear. I was put on high dose steroids. Also having tinnitus (mostly whooshing sound of my own pulse). MRI negative. Blood work negative. Some mild improvement now, after 1 dose steroids.

VAERS ID: 931399 (history)
Form: Version 2.0
Age: 24.0
Sex: Female

Location: Indiana
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood test, Chills, Dyspnoea, Headache, Hypertension, Pyrexia, Tachycardia
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Hypertension (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Depression
Allergies: Toradol: hives Vancomycin: itch
Diagnostic Lab Data: Blood labs 01/08-09/2021
CDC Split Type:

Write-up: Fever up to 102.9, chills, headache, hypertension, tachycardia, dyspnea.

VAERS ID: 931417 (history)
Form: Version 2.0
Age: 60.0
Sex: Male
Location: Florida
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1283 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Activated partial thromboplastin time prolonged, Arthralgia, Blood chloride decreased, Blood glucose normal, Blood potassium decreased, Blood sodium decreased, Cardiac ventriculogram left, Catheterisation cardiac abnormal, Chest pain, Dyspnoea, Echocardiogram normal, Ejection fraction decreased, Electrocardiogram, Electrocardiogram ST segment elevation, Hypokinesia, Myocardial infarction, N-terminal prohormone brain natriuretic peptide increased, Neck pain, Pain, Prothrombin time prolonged, Pyrexia, SARS-CoV-2 test negative, Stent placement, Troponin T increased
SMQs:, Cardiac failure (narrow), Liver-related coagulation and bleeding disturbances (narrow), Anaphylactic reaction (broad), Haemorrhage laboratory terms (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Embolic and thrombotic events, arterial (narrow), Parkinson-like events (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Hyponatraemia/SIADH (narrow), Cardiomyopathy (narrow), Hypotonic-hyporesponsive episode (broad), Chronic kidney disease (broad), Arthritis (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypokalaemia (narrow), COVID-19 (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: zinc sulfate omeprazole metoprolol levothyroxine sodium HCTZ echinacea purpurea extract cholecalciferol
Current Illness: malignant melanoma HTN hypothyroidism GERD
Preexisting Conditions: malignant melanoma
Allergies: codene lactose
Diagnostic Lab Data: portable echo in ER: no effusion, inferior wall poorly visualized EKG: suggestive of Inferior Wall STEMI cardiac cath: RCA-Pd 100% at origin with TIMI 0 flow (infarct related artery) LV angio: inferior and inf-apical hypokinesis, EF approx 45% PCI (drug eluting stent placed to RCA-Pd) troponin T 4.250 NT Pro BNP 2231 SARS CoV-2 (agent for COVID -19) not detected by PCR glucose 106 Na 127 K 3.3 Cl 93 PTT 36.4 PT 13.7
CDC Split Type:

Write-up: Myocardial Infarction: patient began to complain of severe chest pain 3 hours after the vaccine was given .. Vaccine NDC # 59267-1000-1. 0.3 ml given by RN. Patient called his PCP: "... I had very bad chest and shoulder pains, neck pains and slight fever from 9 pm until early this morning (Jan 8). My blood pressure was 155/95 mmHg. Should I see you today? Still feel sore all upper body. Above message received at 0720 am (Jan 8) and the patient was called back at 0757 am (Jan 8): patient was told that many of the side effects above were related to the vaccine but the chest pain was worrisome and the provider requested the

patient go to the emergency room. Patient understood the importance to seek medical attention..... Emergency Room notes: seen by MD on Jan 9. Note at 0749: patient complained of chest pain on/off since received COVID vaccine on Jan 7. Pain was substernal and radiated to the left shoulder, assoc with some SOB. EKG obtained and revealed ST segment elevation and a "cardiac alert" was called.

VAERS ID: 931432 (history)
Form: Version 2.0
Age: 34.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Computerised tomogram head normal, Electroencephalogram normal, Facial paralysis, Fatigue, Hypoaesthesia oral, Laboratory test, Magnetic resonance imaging normal, Muscle twitching, Paraesthesia, Scan with contrast, Ultrasound Doppler normal
SMQs:, Peripheral neuropathy (broad), Dyskinesia (broad), Dystonia (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Hearing impairment (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zoloft 50 mg Clonazepam 0.5mg
Current Illness: none
Preexisting Conditions: none
Allergies: none
Diagnostic Lab Data: all done Jan 6th CT scan - normal EEG- normal MRI with and without contrast - normal doppler of both carotid arteries- normal echo - not performed
CDC Split Type:

Write-up: At around 11:45pm tingling started in my left eyebrow. I thought there was something in my eyebrow and after time of wiping it a few times I went to look into the mirror

and realized that nothing was there. about and hour later I got a drink and a snack to eat and I realized I had a numb sensation in the left corner of my lip. As a nurse I went though the signs of a stroke with 2 other nurses I called. Everything was normal other than that sensation. Thinking I was just overly tired I went to bed. When I woke up the next day I felt okay until I went to drink some coffee and the numb sensation in my corner lip was still there. Now I was concerned and called employee health and was instructed to go to the ER to role out a stoke. I followed employee health"s instructions and went to the ER. I was diagnosed with facial paresthesia and discharged with instructions to come back if symptoms got worse. Symptoms persisted all day and around 7 I called my mom on video chat to show a visual facial twitch on the left side of my face. right after I got off the phone with her the left side of my face drooped and my fiance immediately drove me to the closest ER. I was seen immediately and they decided to admit me. The did labs and head CT. They admitted me to labor and delivery because there were no other rooms. The next day I was seen by a neurologist and many doctors.

VAERS ID: 931484 (history)
Form: Version 2.0
Age: 78.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-01-08
Onset: 2021-01-08
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1686 / 1	AR / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood glucose normal, Fatigue, Loss of consciousness, Pain in extremity, Unresponsive to stimuli
SMQs:, Torsade de pointes/QT prolongation (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient was unresponsive in her room during the night, had gotten the vaccine this morning, 911 called. Had right arm pain and loss of consciousness. EMS got 180/104 BP and blood glucose was 122. Was transported to hospital. Returned to the facility the next day with no complications, was just fatigued.

VAERS ID: 931507 (history)
Form: Version 2.0
Age: 62.0
Sex: Female
Location: New York
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	012L 20A / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Abdominal pain upper, Asthenia, Cardiac stress test, Cardiovascular evaluation, Chest X-ray, Chest discomfort, Dizziness, Electrocardiogram ST segment elevation, Gait inability, Malaise, Myocardial necrosis marker, Nausea, Scan myocardial perfusion, Speech disorder
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Myocardial infarction (broad), Anticholinergic syndrome (broad), Dementia (broad), Dystonia (broad), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Lord attain Trazadone
Current Illness: No
Preexisting Conditions: No

Allergies: No
Diagnostic Lab Data: EKG, chest X-ray, cardiac markers , stress test, nuclear stress scan.
CDC Split Type:

Write-up: 8 hours after vaccine I experienced stomach pain and nausea. I then became very ill and extremely weak. I was laying on floor. Very difficult to talk and very dizzy. Unable to walk. Called 911 and went to ER. Had chest pressure and cardia work up was done. Dx wi ST elevation.

VAERS ID: 931558 (history)
Form: Version 2.0
Age: 47.0
Sex: Female
Location: Arkansas
Vaccinated: 2020-12-22
Onset: 2020-12-30
Days after vaccination: 8
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	- / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Cerebrovascular accident, Computerised tomogram, Hypercoagulation, Injection site pruritus, Injection site swelling, Injection site warmth, Laboratory test, Magnetic resonance imaging brain, Magnetic resonance imaging neck, Scan with contrast
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: CT with contrast, MRI and MRA of head, MRI neck, expensive labwork
CDC Split Type:

Write-up: 7 day after site itching, hot swelling. Unsure if related 9 day after suffered CVA and have hyper coagulation

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VAERS ID: 931658 (history)
Form: Version 2.0
Age: 52.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2021-01-09
Onset: 2021-01-09
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Chest pain, Dizziness, Feeling hot, Immediate post-injection reaction, Retching
SMQs:, Anticholinergic syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: gummy multivitamin
Current Illness: none
Preexisting Conditions: none
Allergies: Demerol, PCN, onions
Diagnostic Lab Data:
CDC Split Type:

Write-up: after receiving vaccine patient immediately felt warm, dizzy and started dry heaving. we dosed Zofran 4mg, gave one dose epi-pen, and 50mg Benadryl. patient still complained of chest pain and was dry heaving. she was then transported by ems to the hospital at 4pm.

VAERS ID: 931662 (history)

Form: Version 2.0
Age: 49.0
Sex: Female
Location: North Carolina
Vaccinated: 2020-12-19
Onset: 2020-12-19
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	ERS730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Appendicectomy, Computerised tomogram, Full blood count, Metabolic function test, Post procedural haemorrhage, Surgical procedure repeated, Vomiting
SMQs:, Acute pancreatitis (broad), Haemorrhage terms (excl laboratory terms) (narrow), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zyrtec, vitamin c, singular,, lisinopril, xolair
Current Illness: Asthma, hypertension
Preexisting Conditions: Asthma, hypertension
Allergies: Lovenox, latex, pollen, mold, cat dander
Diagnostic Lab Data: CT scan, CBC, chem 7 on 12/20/2020
CDC Split Type:

Write-up: Received vaccine 12/19/2020 at hospital around lunch time. Severe abd pain started 10 hours after vaccine administration with vomiting. I went to emergency room next morning Emergency appendectomy 12/20/2020. Had bleeding after surgery. Second surgery 12/21/2020

VAERS ID: 932059 (history)
Form: Version 2.0
Age: 49.0
Sex: Male
Location: Louisiana
Vaccinated: 2021-01-08

Onset: 2021-01-09
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Asthenia, Balance disorder, Diarrhoea, Dizziness, Electrocardiogram abnormal, Fatigue, Feeling abnormal, Full blood count normal, Gait disturbance, Influenza virus test, Influenza virus test negative, Injection site pain, Metabolic function test, Metabolic function test normal, Nausea, Pain, Pallor, Presyncope, Pyrexia, SARS-CoV-2 test negative, Sensory disturbance, Troponin normal, Ventricular extrasystoles, Vomiting
SMQs:, Acute pancreatitis (broad), Peripheral neuropathy (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Ventricular tachyarrhythmias (narrow), Dementia (broad), Pseudomembranous colitis (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Extravasation events (injections, infusions and implants) (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Hypokalaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: amlodipine, losartan, Naprosyn, metoprolol succinate, adipex-p, Topamax, testosterone, zinc gluconate, anastrozole, omeprazole
Current Illness: none
Preexisting Conditions: hypertension
Allergies: KNDA
Diagnostic Lab Data: 01/09/21: CBC- normal CMP- normal Rapid Covid and Flu - negative Troponin- <0.02 EKG- NSR with occasional PVC''s
CDC Split Type:

Write-up: Dizziness started within 30 minutes after injection on 01/08/21 and felt "off". On 01/09/21 approx. 0800 Patient began feeling body aches, fevers, injection site pain, increased dizziness, and nausea. At approximately 1pm patient began vomiting and having diarrhea. Symptoms worsened over the next couple hours to where patient was unable to walk without stumbling. Wife witnessed patient becoming very pale and almost pass out at approx. 5:30pm. Patient states he feels like he''s in slow motion. Patient is unable to maintain balance when walking and reports increasing fatigue and weakness.

VAERS ID: 956978 (history)

Form: Version 2.0
Age: 80.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-22
Onset: 2020-12-29
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Angiogram pulmonary abnormal, Breath sounds abnormal, Chest X-ray abnormal, Cough, Dyspnoea, Pain, Pneumonia, Rhinorrhoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: pneumonia Narrative: On 11/9/20, Patient had a presumptive positive COVID (COBAS) screen as part of routine CLC screening and then on 11/13/20, he had a repeat COVID (CEPHID) that was negative. Then on 12/22/20, he received his first COVID vaccine. On 12/26/20, he began to have c/o hurting all over. Noted history of aspiration and COPD. On 12/29/20, he began to have coughing, increased shortness of breath and runny nose with course breath sounds in his bilateral lower lobes. A chest xray was done and he was initiated on oral azithromycin and cefepime for a bilateral pneumonia. On 1/3/21, he continued to decline with increasing shortness of breath and was subsequently transferred to acute care medicine. All COVID tests have been negative since the presumptive positive on 11/9/20. He did have a CTA that ruled out PE but did show bilateral pneumonia. His antibiotics have been changed to meropenem, vancomycin and IV azithromycin. He remains on acute care at time and has not required ICU care.

VAERS ID: 956991 (history)

Form: Version 2.0
Age: 77.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-28
Onset: 2021-01-04
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Blood urea nitrogen/creatinine ratio increased, Hypovolaemia, SARS-CoV-2 test positive
SMQs:, Acute renal failure (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: COVID positive Narrative: Patient is a resident and received his first COVID vaccine on 12/28/20. On 1/4/21, he had a COVID routine screen done that returned positive on 1/6/21. Per notes, he was asymptomatic at the time; however for isolation purposes, he was transferred to acute care medicine services. On 1/7/21, he was noted to have a slight increase in BUN/creatinine ratio thought to be due to volume depletion and has been ordered IV fluids. He still remains free of any respiratory symptoms at this time.

VAERS ID: 957037 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: Unknown

Vaccinated: 2020-12-23
Onset: 2020-12-26
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Diarrhoea, Nausea, Vomiting
SMQs:, Acute pancreatitis (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Diarrhea & NauseaVomiting Narrative:

VAERS ID: 964641 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Unknown
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Impaired driving ability, Seizure
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis

(broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Narrative: Temporary restriction on driving until further evaluation due to symptoms of seizures.

VAERS ID: 1395602 (history)
Form: Version 2.0
Age: 49.0
Sex: Male
Location: Unknown
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Chills, Computerised tomogram, Dizziness, Ear pain, Fatigue, Headache, Hypoaesthesia, Myalgia, Nausea, Paraesthesia, Tinnitus, Vision blurred, Vomiting
SMQs:, Rhabdomyolysis/myopathy (broad), Acute pancreatitis (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Glaucoma (broad), Lens disorders (broad), Eosinophilic pneumonia (broad), Retinal disorders (broad), Hearing impairment (narrow), Vestibular disorders (broad), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Dizziness, Headache, Myalgia, NauseaVomiting, Fatigue, chills, Right ear pain, dizziness, fuzzy vision, Right sided numbness Narrative: Employee reported fatigue, chills, myalgia, severe right ear pain (drilling like sensation), dizzines, nausea, vomiting and right sided numbness / tingling. CT completed - suspected intracranial bleeding. Transferred to local ED for care and management.

VAERS ID: 932091 (history)
Form: Version 2.0
Age: 50.0
Sex: Male
Location: Florida
Vaccinated: 2020-12-16
Onset: 2021-01-05
Days after vaccination: 20
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / IM

Administered by: Work **Purchased by:** ?
Symptoms: Abdominal pain, Computerised tomogram abdomen abnormal, Mesenteric vein thrombosis
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Embolic and thrombotic events, venous (narrow), Thrombophlebitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Ischaemic colitis (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 5 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: NO
Current Illness: NO
Preexisting Conditions: None
Allergies: Simvastatin
Diagnostic Lab Data:
CDC Split Type:

Write-up: Started severe belly pain and went to Emergency room and diagnosed with mesenteric vein thrombosis after the CT scan of the abdomen, treated with heparin drip, antibiotic and discharged with anticoagulant pills(Eliquis). I am not sure that it is because of the vaccine my doctors are also not sure about it, but I am sure that I am a healthy person without any health issues . I am working as registered nurse, our unit is for covid-19 patient"s since march 2020 and I had covid -19 on August month and recovered after 3 weeks.

VAERS ID: 932145 (history)
Form: Version 2.0
Age: 96.0
Sex: Female
Location: North Carolina
Vaccinated: 2021-01-08
Onset: 2021-01-08
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3248 / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram cerebral, Aphasia, Arteriogram carotid, Cerebral artery occlusion, Computerised tomogram head, Facial paresis, Hypotonia, Ischaemic stroke, Monoparesis, NIH stroke scale abnormal
SMQs:, Peripheral neuropathy (broad), Ischaemic central nervous system vascular conditions (narrow), Dementia (broad), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Hypertension, hypothyroidism
Preexisting Conditions: Hypertension, hypothyroidism
Allergies: None
Diagnostic Lab Data: Complete stroke workup with noncontrast CT head and CT angiogram of the head and neck.
CDC Split Type:

Write-up: Patient came into the emergency department on 1/8/21 with an acute ischemic stroke with complete occlusion of her left MCA. She had acute and complete flaccid paresis of her right face, arm, and leg, complete aphasia, and neglect of the right side of her body. NIHSS of 27. Onset of deficit was between 6:30pm-7:10pm. She recieved her 1st COVID-19 vaccine dose that morning at 10:31am.

VAERS ID: 932366 (history)
Form: Version 2.0
Age: 38.0
Sex: Female
Location: Virginia
Vaccinated: 2021-01-09
Onset: 2021-01-09
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3246 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram cerebral, Arteriogram carotid, Chest X-ray, Computerised tomogram head, Facial paresis, Full blood count, Hypoaesthesia, International normalised ratio, Magnetic resonance imaging, Metabolic function test, Muscular weakness, Prothrombin time, Visual field defect
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Optic nerve disorders (broad), Retinal disorders (broad), Immune-mediated/autoimmune disorders (broad), Sexual dysfunction (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes

Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: with past 24 hours prior to vxn pt reports taking: One a day women''s vitamin, Benadryl, Allegra
Current Illness: none
Preexisting Conditions: asthma, cold sores
Allergies: Pineapple
Diagnostic Lab Data: 1/9 - CT of Head, CTA head and neck, CBC, PT/INR, BMP, Chest xray 1/10 - Repeat CT of head, MRI,
CDC Split Type:

Write-up: The patient presented with left eye peripheral visual loss, left upper and lower extremity and facial numbness sensation and weakness. This started 1 hour after receiving COVID-19 vaccine at her place of employment. Pt was brought to CRMC via EMS.

VAERS ID: 932372 (history)
Form: Version 2.0
Age: 93.0
Sex: Male
Location: West Virginia
Vaccinated: 2021-01-02
Onset: 2021-01-02
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / IM

Administered by: Public **Purchased by:** ?
Symptoms: Asthenia, Cough, Gait inability, Malaise, Pyrexia, SARS-CoV-2 test positive
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dystonia (broad), Guillain-Barre syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Amlodipine, NPH Insulin, Levothyroxine, Simvastatin
Current Illness:
Preexisting Conditions:
Allergies: No known allergies

Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient received the Moderna Vaccine 1/2/21 at his VA Clinic. He received the vaccine that morning and by the evening he was not feeling well. He developed cough, weakness and fever and now is unable to ambulate. He has since been hospitalized and has tested positive for COVID-19.

VAERS ID: 932480 (history)
Form: Version 2.0
Age: 69.0
Sex: Male
Location: Minnesota
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood test normal, Confusional state, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Covid 19 infection diagnosed 5weeks prior
Preexisting Conditions: COPD, CAD, HTN, demenita
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: pt received Moderna vaccine. next day he had high fevers up to 103, confusion. admitted to hospital. infectious work up negative. improved off antibiotics.

VAERS ID: 932515 (history)

Form: Version 2.0
Age: 73.0
Sex: Male
Location: Ohio
Vaccinated: 2020-12-29
Onset: 2020-12-31
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	E10140 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: COVID-19 pneumonia, Chromaturia, Computerised tomogram, Dyspnoea, Jaundice, Laboratory test, Malaise, Ocular icterus, Yellow skin
SMQs:, Rhabdomyolysis/myopathy (broad), Cholestasis and jaundice of hepatic origin (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Acute central respiratory depression (broad), Biliary system related investigations, signs and symptoms (narrow), Biliary tract disorders (narrow), Pulmonary hypertension (broad), Cardiomyopathy (broad), Conjunctival disorders (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Asprin, cyanacabalmin, dulpxetine, lasix, isosorbide, Slisinop
Current Illness: COPD, PTSD, CAD, BPH, Post COVID
Preexisting Conditions: CAD, COPD, PTSD, BPH
Allergies: PCN, Lactose intolerant
Diagnostic Lab Data: CT Scan, Lab work,
CDC Split Type:

Write-up: Resident appeared to be jaundice with yellow skin and eyes. Resident also complained of not feeling well. Urine was dark yellow. Resident short of breath. Resident was admitted to hospital and diagnosed with post-covid pneumonia.

VAERS ID: 932623 (history)
Form: Version 2.0
Age: 78.0

Sex: Male
Location: Illinois
Vaccinated: 2021-01-09
Onset: 2021-01-10
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1283 / 2	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Angiogram, Basilar artery occlusion, Computerised tomogram, Ischaemic stroke
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: amlodipine, atorvastatin, esomeprazole, metoprolol, omega-3, ticagrelor, cyanocobalamin
Current Illness:
Preexisting Conditions: Hypertension, hyperlipidemia, coronary artery disease (s/p angioplasty)
Allergies: aspirin
Diagnostic Lab Data: CT/CTA
CDC Split Type:

Write-up: Acute ischemic stroke, basilar occlusion

VAERS ID: 932668 (history)
Form: Version 2.0
Age: 55.0
Sex: Male
Location: Illinois
Vaccinated: 2021-01-07
Onset: 2021-01-09
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Dyspnoea
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Senna 8.6mg 2 tabs po Vit. D 1000 ui 2 tabs po Lactulose 30 cc BID
Current Illness: none
Preexisting Conditions: 318.2 profound intellectual disability gingivitis 758.0 down syndrome 272.4 hyperlipidemia 564.00 chronic constipation vitamin D insufficiensy
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: SOB

VAERS ID: 932696 (history)
Form: Version 2.0
Age: 26.0
Sex: Male
Location: Illinois
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	AR / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Decreased appetite, Dehydration, Diarrhoea, Nausea, Tachycardia, Vomiting
SMQs:, Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow), Dehydration (narrow)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Keppra,colase,miralax, erythromycin, acidophilus
Current Illness: no
Preexisting Conditions: CP, seizure disorder
Allergies: Gluten, lactose intolerance
Diagnostic Lab Data: 1/10/21 sent to er Dx dehydration
CDC Split Type:

Write-up: Nausea/vomiting and diarrhea loss of appetite tachicardia sent to er for hydration.

VAERS ID: 932801 (history)
Form: Version 2.0
Age: 32.0
Sex: Male
Location: Illinois
Vaccinated: 2020-12-17
Onset: 2020-12-19
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Abdominal pain upper, Condition aggravated, Gallbladder disorder, Gallbladder operation
SMQs:, Acute pancreatitis (broad), Gallbladder related disorders (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: Unknown
Current Illness:
Preexisting Conditions: Hx of gallbladder disease
Allergies: Unknown
Diagnostic Lab Data:
CDC Split Type:

Write-up: Two days following the first COVID vaccination he developed epigastric pain and was evaluated in the ER and was admitted for gallbladder surgery. He has a 6 year history of gallbladder disease but was not experiencing symptoms until after the COVID vaccination.

VAERS ID: 932915 (history)
Form: Version 2.0
Age: 25.0
Sex: Female
Location: Washington
Vaccinated: 2020-12-29
Onset: 2021-01-08
Days after vaccination: 10
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Contusion, Haematocrit normal, Haemoglobin normal, Mean cell haemoglobin concentration normal, Mean cell haemoglobin normal, Mean cell volume normal, Mouth haemorrhage, Platelet count decreased, Red blood cell count normal, Red cell distribution width normal, Thrombocytopenia, White blood cell count normal
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Accidents and injuries (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zoloft (sertraline), vitamin D, cetirizine (zyrtec)
Current Illness: None
Preexisting Conditions: Anxiety, history of anti-thyroglobulin antibodies
Allergies: None
Diagnostic Lab Data: CBC 1/8/21 - WBCs 8.65, RBCs 4.47, Hg 13.7, Hct 40, MCV 89, MCH 30.6, MCHC 34.3, Plts 1** (d) C, RDW 11.8%.

CDC Split Type:

Write-up: Severe thrombocytopenia (plts 3k/uL), oral mucosal bleeding, bruising

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VAERS ID: 933025 (history)
Form: Version 2.0
Age: 43.0
Sex: Female
Location: Arizona
Vaccinated: 2020-12-21
Onset: 2020-12-28
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Angiogram cerebral normal, Anion gap, Arteriogram carotid normal, Blood calcium normal, Blood chloride normal, Blood creatinine normal, Blood glucose normal, Blood potassium decreased, Blood sodium normal, Carbon dioxide decreased, Computerised tomogram head abnormal, Encephalitis, Glomerular filtration rate normal, Glycosylated haemoglobin normal, Headache, Hypokalaemia, Inflammation, Ketonuria, Magnetic resonance imaging brain normal, Oral herpes, Platelet count normal, Posterior reversible encephalopathy syndrome, Urine ketone body present, White blood cell count normal
SMQs:, Hyperglycaemia/new onset diabetes mellitus (narrow), Oropharyngeal infections (narrow), Noninfectious encephalitis (narrow), Noninfectious encephalopathy/delirium (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypokalaemia (narrow), Opportunistic infections (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: methimazole 2.5 mg daily, MVI, D3, B12
Current Illness: none
Preexisting Conditions: hyperthyroidism
Allergies: None
Diagnostic Lab Data: CBC: normal WBC, no anemia, normal platelet BMP: glucose 115, eGFR 96 Cr 0.76, Na 136, K 3.3, Cl 101 CO2 22, AG 13, Ca 9.2 urine + ketones (small) CT head: IMPRESSION: 1. No evidence of acute parenchymal hemorrhage 2. Patchy areas of hypodensity noted within the occipital lobes bilaterally greater on the left than the right. Findings favored to represent underlying posterior reversible encephalopathy syndrome,

further evaluation with MRI of the brain is warranted. CT angio head/neck Impression: 1. No evidence of significant stenosis, vascular occlusion, arterial dissection extracranial vasculature of the neck 2. No evidence of significant stenosis, vascular occlusion, arterial aneurysm the arteries of the circle of Willis MRI brain w/o contrast IMPRESSION: No evidence of an acute intracranial process. Essentially unremarkable examination of the brain without distinct findings to indicate PRESS syndrome. 12/29 labs indicated resolution of hypokalemia and ketonuria. Hgb A1c 5.5%.
CDC Split Type:

Write-up: Sudden, severe worst headache of her life, coupled with onset of oral herpetic lesions and inflammation to unrelated body part (belly button piercing) occurred one week after vaccination received. She presented to ED 12/28 noted to have hypokalemia and head CT showed mild occipital encephalitis, admitted overnight for obs subsequent brain MRI was normal. She was seen in my clinic 2 days later (1/4/21) and was started on 3 day course of Decadron, topical acyclovir for herpetic lesion. As she is a nurse, I kept her off work until resolution of symptoms. Seen again on 1/8/21, headache resolved. She did discuss CT and MRI results with neurologist. He was not convinced this was vaccine related. She is having 2nd dose of vaccine on 1/11/21.

VAERS ID: 951288 (history)
Form: Version 2.0
Age: 69.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 2021-01-06
Submitted: 0000-00-00
Entered: 2021-01-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Blood potassium decreased, Dyspnoea, Respiratory depression
SMQs:, Anaphylactic reaction (broad), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Cardiomyopathy (broad), Respiratory failure (narrow), Hypokalaemia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: RespDepression found to have low potassium Narrative: Patient reports experiencing shortness of breath 24 hours after the vaccine was administered, went to ER and admitted for one night. Informed her supervisor she had low potassium.

VAERS ID: 933299 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Encephalitis, Intensive care
SMQs:, Noninfectious encephalitis (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Drug allergy
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021007424

Write-up: encephalitis; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received BNT162B2(PFIZER-BIONTECH COVID-19 VACCINE) , via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The patient was allergic to UTI(urinary tract infection) infection medication (i.e. sulfamethoxazole, trimethoprim etc.). Concomitant medications were not reported. The patient got encephalitis and was put in the ICU(intensive care unit) after getting

vaccinated, led to hospitalization. Outcome of event was unknown. Information about lot/batch number has been requested.

VAERS ID: 933369 (history)
Form: Version 2.0
Age: 56.0
Sex: Female
Location: Minnesota
Vaccinated: 2021-01-08
Onset: 2021-01-08
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction
SMQs:, Anaphylactic reaction (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypersensitivity (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Albuteral, Loratadine, Fmotidine, Prednisone, Levothyroxine, Epipen (prn)
Current Illness: None
Preexisting Conditions: Migraine headaches, History of papillary thyroid carcinoma, Latex allergy
Allergies: Latex
Diagnostic Lab Data:
CDC Split Type:

Write-up: Anaphylactic reaction

VAERS ID: 933739 (history)
Form: Version 2.0
Age: 54.0

Sex: Female
Location: Ohio
Vaccinated: 2021-01-08
Onset: 2021-01-09
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Brain death, Bronchial secretion retention, Cardio-respiratory arrest, Death, Dyspnoea, Electroencephalogram abnormal, Intensive care, Mechanical ventilation, Resuscitation, SARS-CoV-2 test negative, Withdrawal of life support
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (narrow), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Cardiomyopathy (broad), Respiratory failure (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-10
Days after onset: 1
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Aspirin, Diazepam, Doxepin HCL, Duloxetine, Juven POW, Lamotri Levothyroxin, Loratadine, Melatonin, Mucus Relief, Olanzapine, Prazosin HCL, Pregabalin,
Current Illness: Recent g-tube placement, Several hospitalizations over the pas few months due to low Oxygen Levels.
Preexisting Conditions: Major depression, borderline personality disorder, Cerebral Palsy, History of dissected left carotid artery
Allergies: Penicillin
Diagnostic Lab Data: COVID-19 test administered - 1/9/2021 - Negative 2 EEGs were performed on 1/9/2021 and both indicated she had not brain activity
CDC Split Type:

Write-up: Staff member checked on her at 3am and patient stated that she felt like she couldn"t breathe. 911 was called and taken to the hospital. While in the ambulance, patient coded. Patient was given CPR and "brought back". Once at the hospital, patient was placed on a ventilator and efforts were made to contact the guardian for end of life decisions. Two EEGs were given to determine that patient had no brain activity. Guardian, made the decision to end all life saving measures. Patient was taken off the ventilator on 1/9/2021 and passed

away at 1:30am on 1/10/2021. The initial indication from the ICU doctor was the patient had a mucus plug that she couldn''t clear.

VAERS ID: 933751 (history)
Form: Version 2.0
Age: 58.0
Sex: Male
Location: Pennsylvania
Vaccinated: 2021-01-11
Onset: 2021-01-11
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / 1	LA / IM

Administered by: Military **Purchased by:** ?
Symptoms: Dyspnoea, Malaise, Oxygen saturation normal, Throat tightness
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Advair 500/50, Spriva 2.5mcg, Asmanex 220mcg, Metformin 100mcg, Nexium 20mg, Montelekast, Baby asprin 81 mg, Lovastatin, Nosonex, Pro Air, Duo Neb, Epi Pen 0.3 mg Busporine 150 Mg, Theophylin 400mg.
Current Illness:
Preexisting Conditions:
Allergies: Shell Fish, Cologne, Perfume, Hand sanitizer, penicillin, Zpac, coconut, peanuts
Diagnostic Lab Data:
CDC Split Type:

Write-up: Aprox 5 minutes after vaccine was given, patient stated he was not feeling well, asked for water and stated he was having trouble breathing. Was then assisted to clinic treatment room, vitals were taken: 08:45 B/P 171/110, HR 84, R 28 O2 97. EMS was activated. At 0900 158/94 HR 84 R 24 O2 97. Patient then voice he was having tightness in his throat. Epi Pen 0.3mg administered at 0900. At 0904 EMS arrived and patient taken to the nearest hospital.

VAERS ID: 933755 (history)

Form: Version 2.0
Age: 26.0
Sex: Female
Location: Maryland
Vaccinated: 2021-01-03
Onset: 2021-01-04
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Hypoaesthesia, Monoplegia, Paraesthesia, Seizure
SMQs:, Peripheral neuropathy (broad), Systemic lupus erythematosus (broad), Convulsions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Numbness and tingling bil upper extremities, seizure, temporary paralysis R arm. Started day after vaccine given, was observed overnight in hospital.

VAERS ID: 933792 (history)
Form: Version 2.0
Age: 33.0
Sex: Female
Location: Maine

Vaccinated: 2020-12-23
Onset: 2021-01-01
Days after vaccination: 9
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Neurological symptom
SMQs:

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: None
Diagnostic Lab Data:
CDC Split Type:

Write-up: Manager was notified that the employee had suffered a stoke-like "serious medical event" hours after the end of her shift, while at home.

VAERS ID: 933805 (history)
Form: Version 2.0
Age: 66.0
Sex: Female
Location: Alabama
Vaccinated: 2020-12-16
Onset: 2020-12-21
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	- / IM

Administered by: Work **Purchased by:** ?
Symptoms: Arthralgia, Body temperature increased, Headache, Neck pain, Odynophagia

SMQs:, Neuroleptic malignant syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Arthritis (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient had vaccine in Left arm. That same night patient had temp of 100.1 and the right neck at base of head to shoulder began to hurt patient was unable to swallow without pain in the next few days. Patient went to ER and was hospitalized for 2 days treated with IV steroids and 2 antibiotics (clindamycin and acyclovir). Patients symptoms resolved and patient was discharged without additional issues. The admitting physician was unable to identify the cause of these symptoms, but the vaccine could not be ruled out.

VAERS ID: 933913 (history)
Form: Version 2.0
Age: 65.0
Sex: Female
Location: Indiana
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Fatigue, Myalgia, Respiratory rate increased
SMQs:, Rhabdomyolysis/myopathy (broad), Eosinophilic pneumonia (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Prescription 03/19/2019 - Open Ended metoprolol succinate tablet extended release 24 hr; 50 mg; amt: 50 mg; oral Special Instructions: Hold for SBP <100 or HR <60 Once A Day; Upon Rising Prescription 07/12/2019 - Open Ended Ampyra (dalfamp
Current Illness:
Preexisting Conditions: Neuromuscular dysfunction of bladder, unspecified 02/26/2018 R13.13 Dysphagia, pharyngeal phase 03/06/2020 G35 Multiple sclerosis 02/26/2018 K58.9 Irritable bowel syndrome without diarrhea 02/26/2018 I50.9 Heart failure, unspecified 02/26/2018 K21.9 Gastro-esophageal reflux disease without esophagitis 02/26/2018 F33.1 Major depressive disorder, recurrent, moderate 03/18/2019 I10 Essential (primary) hypertension 02/26/2018 F41.1 Generalized anxiety disorder 02/26/2018 F39 Unspecified mood [affective] disorder 02/26/2018 H04.129 Dry eye syndrome of unspecified lacrimal gland 02/26/2018 G47.8 Other sleep disorders 04/04/2019 E87.5 Hyperkalemia 11/13/2018 M62.838 Other muscle spasm 02/26/2018 J30.9 Allergic rhinitis, unspecified 11/25/2019 R26.2 Difficulty in walking, not elsewhere classified 09/14/2018 E53.8 Deficiency of other specified B group vitamins 06/28/2018 E55.9 Vitamin D deficiency, unspecified 02/26/2018 E56.8 Deficiency of other vitamins 02/26/2018 R42 Dizziness and giddiness 02/26/2018 R52 Pain, unspecified 02/26/2018 R13.13 Dysphagia, pharyngeal phase (History of) 05/10/2019 R53.1 Weakness 09/14/2018 R26.81 Unsteadiness on feet 03/06/2020 R26.89 Other abnormalities of gait and mobility 03/06/2020 M62.81 Muscle weakness (generalized) 02/26/2018 Z51.89 Encounter for other specified aftercare 10/03/2018 Z86.16 Personal history of COVID-19 Note:June 2020 01/07/2021 R26.2 Difficulty in walking, not elsewhere classified 07/17/2020 R26.81 Unsteadiness on feet 07/17/2020 R26.89 Other abnormalities of gait and mobility 07/17/2020 G35 Multiple sclerosis
Allergies: gabapentin, guaifenesin, Penicillins (PCN)
Diagnostic Lab Data:
CDC Split Type:

Write-up: Upon assessment resident noted to have increased respirations, lungs CTA. Resident c/o increased fatigue and muscle aches. VS 202/180, 118, 22, 97.1, 96%.

VAERS ID: 933935 (history)
Form: Version 2.0
Age: 43.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-01-02
Onset: 2021-01-10
Days after vaccination: 8
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	AR / IM

Administered by: Private **Purchased by:** ?

Symptoms: Haemoglobin normal, Immune thrombocytopenia, Platelet count decreased, Thrombocytopenia, White blood cell count normal
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Omeprazole
Current Illness: None
Preexisting Conditions: GERD
Allergies: NKDA
Diagnostic Lab Data: CBC showed platelet count of 2,000. WBC and Hgb normal
CDC Split Type:

Write-up: Sever thrombocytopenia (platelet count 2,000) 8 days following Moderna COVID vaccine. Clinically suspicious for ITP.

VAERS ID: 933950 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: D.C.
Vaccinated: 2021-01-05
Onset: 2021-01-07
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Eye swelling, Fatigue, Peripheral swelling, Swelling face
SMQs:, Cardiac failure (broad), Anaphylactic reaction (broad), Angioedema (narrow), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: crestor nexium trazodone zanaflex synthroid
Current Illness:
Preexisting Conditions: Ulcerative Colitis Rheumatoid Arthritis
Allergies: mushrooms Humira Pen (site reaction) dextromethorphan erythromycin (nausea & vomitting) sulfaSALAzine (hives)
Diagnostic Lab Data: admitted to the hospital on 1/10/2021
CDC Split Type:

Write-up: Received vaccine on 1/5, began having swelling in bilateral hands and lower extremities on 1/7, along with fatigue. On 1/8 she reports new swelling started in her face (eyes and cheeks). Swelling has not improved today and fatigue has worsened.

VAERS ID: 933965 (history)
Form: Version 2.0
Age: 81.0
Sex: Female
Location: Indiana
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood glucose increased, Confusional state, Discomfort, Myalgia, Pyrexia
SMQs:, Rhabdomyolysis/myopathy (broad), Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Eosinophilic pneumonia (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: allopurinol tablet; 300 mg; amt: 300mg; oral Once A Day; Upon Rising Prescription 04/06/2019 - Open Ended furosemide tablet; 40 mg; amt: 40mg; oral Once A Day; Upon Rising Prescription 04/06/2019 - Open Ended levothyroxinet tablet; 50 mcg;
Current Illness:
Preexisting Conditions: E11.9 Type 2 diabetes mellitus without complications 09/14/2018 J98.4 Other disorders of lung Note:restrictive lung disease 09/14/2018 D50.9 Iron deficiency anemia, unspecified 09/14/2018 E78.5 Hyperlipidemia, unspecified 09/14/2018 I73.9 Peripheral vascular disease, unspecified 09/14/2018 I48.0 Paroxysmal atrial fibrillation 09/14/2018 I13.0 Hypertensive heart and chronic kidney disease with heart failure and stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease 04/06/2019 F41.9 Anxiety disorder, unspecified 09/14/2018 F33.1 Major depressive disorder, recurrent, moderate 02/21/2019 M19.90 Unspecified osteoarthritis, unspecified site 04/06/2019 E03.9 Hypothyroidism, unspecified 09/14/2018 F41.0 Panic disorder [episodic paroxysmal anxiety] 09/14/2018 R44.3 Hallucinations, unspecified 09/14/2018 M10.9 Gout, unspecified 09/14/2018 I50.32 Chronic diastolic (congestive) heart failure 11/01/2018 N18.3 Chronic kidney disease, stage 3 (moderate) 04/06/2019 K30 Functional dyspepsia 09/14/2018 R45.1 Restlessness and agitation 09/14/2018 G47.00 Insomnia, unspecified 09/14/2018 M53.87 Other specified dorsopathies, lumbosacral region Note:DJD 09/14/2018 E04.2 Nontoxic multinodular goiter 09/14/2018 R29.6 Repeated falls 11/01/2018 R26.2 Difficulty in walking, not elsewhere classified 09/14/2018 M62.81 Muscle weakness (generalized) 09/14/2018 R53.1 Weakness 11/01/2018 R26.81 Unsteadiness on feet 09/14/2018 E66.01 Morbid (severe) obesity due to excess calories 09/14/2018 Z87.440 Personal history of urinary (tract) infections 04/07/2019 Z87.891 Personal history of nicotine dependence 04/06/2019 Z87.81 Personal history of (healed) traumatic fracture 04/06/2019 Z51.89 Encounter for other specified aftercare 09/14/2018 R50.9 Fever, unspecified 07/20/2020 R26.89 Other abnormalities of gait and mobility 04/24/2020 L97.811 Non-pressure chronic ulcer of other part of right lower leg limited to breakdown of skin 09/27/2019 M15.0 Primary generalized (osteo)arthritis
Allergies: codeine phosphate (bulk), enalapril maleate, oxycodone-aspirin, Penicillins (PCN)
Diagnostic Lab Data:
CDC Split Type:

Write-up: Found on floor by CNA at 6 am. On assessment by charge nurse, VS: 99.6-94-16 212/105 manual. Noted increased confusion. Temp. 99.6 FBS 114. Resident had been provided tylenol just prior in shift for general discomfort (sore muscles) and low grade temp. On call for Dr notified of all and received order to send to ER.

VAERS ID:	934094 (history)
Form:	Version 2.0
Age:	37.0
Sex:	Female
Location:	Oklahoma
Vaccinated:	2021-01-05
Onset:	2021-01-05
Days after vaccination:	0
Submitted:	0000-00-00

Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Ageusia, Chest discomfort, Cough, Diarrhoea, Dyspnoea, Fatigue, Lip pruritus, Malaise, Oral discomfort, Pruritus, Pyrexia, SARS-CoV-2 test negative
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Taste and smell disorders (narrow), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: humalog,lantus,omepriazole,topamax,prozac,
Current Illness: migraines, asthma
Preexisting Conditions: Diabetes, asthma, migraines
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: Approximately 15 minutes after receiving the vaccine, client started c/o itching to face and lips. Transported to triage area. Where she started c/o burning lips also. vital signs monitored, Benadryl 50mg po given. In a few minutes she reported chest tightness, labored breathing. 1:44pm Epinephrine was given IM. client was on 02 increased to 15 Liters. She admits to decreased chest tightness and itching of face improving. She was taken by ambulance to hospital, where she was admitted overnight for observation. She was given Benadryl and epi X2 more time, once in ambulance and once at the hospital. She denies any problems since then. She did say that approx 3 weeks ago she was a contact to a case of covi

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VAERS ID: 934099 (history)
Form: Version 2.0
Age: 41.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-28

Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	UN / IM

Administered by: Private **Purchased by:** ?
Symptoms: Burning sensation, Diarrhoea, Laboratory test, Nausea, Rash, Rash pruritic, Throat tightness
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (broad), Peripheral neuropathy (broad), Pseudomembranous colitis (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (narrow), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: topomax, melatonin, mirena ring, B12 injection, zyrtec, biotin
Current Illness: none
Preexisting Conditions: anxiety, depression, hyperprolactinemia, migraines, primary hypothyroidism
Allergies: ? shellfish (positive skin test but eats shellfish with "no problem")
Diagnostic Lab Data: just basic labs
CDC Split Type:

Write-up: Patient is a 41 y.o. female who presented to the ED with complaints of a reaction after she took the COVID vaccine. Patient is an RN here and had earlier received COVID-19 vaccination (Moderna) around 0130 this afternoon. Soon after she experienced rash which was burning and itchy on her arms thighs and back. She reported to Occ health and was directed to the ED. While in the ED she experienced throat tightness and nausea. She had one episode of diarrhea. No tongue, lip swelling, SOB or difficulty swallowing. Denies any chest pain, palpitations, pre-syncope/syncope. No vomiting abdominal pain. H/o of allergy to fish mix and dust mite.

VAERS ID: 934156 (history)
Form: Version 2.0
Age: 60.0
Sex: Female
Location: Indiana
Vaccinated: 2021-01-05

Onset: 2021-01-07
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	RA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Chills, Intensive care, Laboratory test, Pain, Unresponsive to stimuli
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Tylenol, Atenolol, LIpitor, Clonidine, mutlivitamin, Lasix, Gabapentin, Losartan, Maalox, Miralax, Zofran, Vitamin D2.
Current Illness: No
Preexisting Conditions: 2019-nCoV acute respiratory disease, Hyperlipidemia, unspecified, Chronic obstructive pulmonary disease, unspecified, Other hereditary and idiopathic neuropathies, Essential (primary) hypertension, Peripheral vascular disease, unspecified, Venous insufficiency (chronic) (peripheral), Other chronic pain, Type 2 diabetes mellitus with hyperglycemia, Dehydration, Nondisplaced oblique fracture of shaft of left fibula, sequela, Vitamin D deficiency, unspecified, Nondisplaced fracture of second metatarsal bone, left foot, subsequent encounter for fracture with routine healing, Nondisplaced fracture of third metatarsal bone, left foot, subsequent encounter for fracture with routine healing, Nondisplaced fracture of fourth metatarsal bone, left foot, subsequent encounter for fracture with routine healing, Hypokalemia, Acute respiratory failure with hypoxia, Muscle weakness (generalized), Difficulty in walking, not elsewhere classified, Cognitive communication deficit, Non-pressure chronic ulcer of other part of left lower leg with fat layer exposed, Repeated falls, Cellulitis of right lower limb, Cellulitis of left lower limb, Other iron deficiency anemias, Other recurrent depressive disorders, Hypothyroidism, unspecified, Chronic venous hypertension (idiopathic) with ulcer of bilateral lower extremity, Diabetes mellitus due to underlying condition with other skin complications, Other constipation, Chronic respiratory failure with hypoxia, Other acute kidney failure, Lymphedema, not elsewhere classified, Generalized edema, Morbid (severe) obesity due to excess calories, Chronic ischemic heart disease, unspecified, Gastro-esophageal reflux disease without esophagitis, Disorder resulting from impaired renal tubular function, unspecified, History of falling, Cellulitis/abscess, leg, Impaired renal function NOS, Hypertension, benign essential
Allergies: Bactrim, Kefles, PCN
Diagnostic Lab Data: All tests done at the Hospital
CDC Split Type:

Write-up: 01/06/21 at 6 pm, body aches, and chills 01/07/21 at 12am T102.2, SPO2 62% on room air. Was sent to ER and returned. 01/08/21 at SPO@ less then 60% on room air, non responsive to verbal tactile stimuli. Responsive to sternal rub only. Was sent to ER and admitted to ICU.

VAERS ID: 934383 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: Maryland
Vaccinated: 2021-01-08
Onset: 2021-01-08
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0142 / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood creatine phosphokinase normal, Blood magnesium normal, Blood thyroid stimulating hormone normal, Borrelia test, Borrelia test negative, Full blood count abnormal, Gait inability, Human chorionic gonadotropin negative, Hypoaesthesia, Hypochromic anaemia, Limb discomfort, Lymphocyte percentage increased, Magnetic resonance imaging brain normal, Magnetic resonance imaging spinal normal, Metabolic function test normal, Mobility decreased, Muscular weakness, Neutrophil percentage decreased, Red blood cell sedimentation rate increased, SARS-CoV-2 test negative, Scan with contrast normal, Treponema test negative, Vitamin B12 normal, Vitamin D decreased
SMQs:, Rhabdomyolysis/myopathy (broad), Agranulocytosis (broad), Haematopoietic leukopenia (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Dystonia (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none
Current Illness: none known
Preexisting Conditions: obesity
Allergies: NKA

Diagnostic Lab Data: 1/10/21: Vitamin D 11 ng/mL, Lyme IgG and IgM negative, Magnesium 2.1 mg/dL, CK 130, HCG negative 1/9/21: RPR nonreactive, SARS COV2 negative 1/8/21: Vitamin B12 540 pg/mL, Sed rate 83 mm/Hr, TSH 4.481 uIU/mL, BMP normal, CBC mild microcytic, hypochromic anemia, lympocytes relative 50.9%, neutrophils relative 42.5% MRI Brain with and without contrast and MRI spinal survey with and without contrast are normal
CDC Split Type:

Write-up: Approximately 10 minutes after receiving the vaccine she started experiencing numbness and inability to move all 4 extremities. No difficulty breathing or swallowing. Benedryl 25 mg was administered with no relief. EMS was called, she was transported to the ED and admitted to the hospital for evaluation. She was given Ativan 0.5 mg IV with some mild improvement in symptoms. She has had gradual improvement in her symptoms, now able to move her arms normally. She has persistent weakness and discomfort in both lower extremities and is unable to ambulate without assistance.

VAERS ID: 934407 (history)
Form: Version 2.0
Age: 98.0
Sex: Female
Location: Connecticut
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL 1284 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Electrocardiogram PR prolongation, Full blood count, Influenza A virus test negative, Influenza B virus test, Influenza virus test negative, Legionella test, Lethargy, Mental status changes, Metabolic function test, Polymerase chain reaction, Pyrexia, Streptococcus test negative, Tachycardia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Conduction defects (narrow), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days

Extended hospital stay? No
Previous Vaccinations:
Other Medications: Tylenol, Artificial tears, Citrical with Vit.D, Ibuprophen, Multivitamin, Prosight, Soothe XP (light mineral oil-mineral oil) eye ointment
Current Illness: Pneumonia, Covid19
Preexisting Conditions: Schizophrenia, unspecified ,Vascular dementia without behavioral disturbance, Venous insufficiency (chronic) (peripheral), Hypo-osmolality and hyponatremia, Unspecified strabismus, Primary osteoarthritis, Unspecified blepharitis unspecified eye, unspecified eyelid
Allergies: Propoxyphene, Unasyn
Diagnostic Lab Data: Blood work-CBC, CMP Done @ ER ECG Done at ER- Result Sinus rhythm, prolonged PR interval. Legionella & Strp Antigen done at ER both were negative. PCR Done at ER for Influenza A&B was negative
CDC Split Type:

Write-up: 1/8/2021 tachycardia Pulse 140, Fever T99.4 then 100.2 Lethargy, Somewhat Altered Mental status

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VAERS ID: 934475 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Michigan
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	RA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Anaphylactic reaction, Dysphonia, Dyspnoea, Peripheral swelling, Speech disorder
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Angioedema (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Dementia (broad), Parkinson-like events (broad), Acute central respiratory depression (broad), Psychosis and psychotic disorders (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ELIQUIS; ; ; CRESTOR; ; SYNTHROID; BENADRYL
Current Illness: Thyroid disorder
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021000598

Write-up: Her voice became raspy, she could hardly talk, she could barely talk; her hand and whole arm swelled up; her hand and whole arm swelled up; Trouble breathing; This is a spontaneous report from a contactable healthcare professional (patient). A 57-year-old female patient received the 1st dose of bnt162b2 (BNT162B2, Manufacturer Pfizer-BioNTech) (lot# EK9231, exp date Apr2021), via an unspecified route of administration in the right arm, on 29Dec2020, at single dose, for COVID-19 immunization and diphenhydramine hydrochloride (BENADRYL), via an unspecified route of administration, at unknown posology, on 29Dec2020 (20 minutes prior to taking bnt162b2), for anaphylactic reaction. Medical history included ongoing thyroid disorder. Concomitant medications included apixaban (ELIQUIS), diltiazem (unknown trade name), losartan (unknown manufacturer), rosuvastatin calcium (CRESTOR), levothyroxine (unknown trade name), ongoing levothyroxine sodium (SYNTHROID) for thyroid disorder and metformin (unknown trade name). Previously the patient received unspecified influenza vaccine (reported as flu shot) for immunisation, on unspecified date, and experienced anaphylactic reaction treated with Benadryl. On 29Dec2020, an hour after getting the vaccine, the patient experienced her voice became raspy, she could hardly talk, she could barely talk with outcome of unknown, her hand and whole arm swelled up with outcome of unknown, trouble breathing with outcome of unknown. The reporter stated that her voice and everything reacted and made her go to the emergency room on 29Dec2020. She reported that had adverse effect more than normal. The event "Her voice became raspy, she could hardly talk, she could barely talk" caused patient"s hospitalization on unknown date. The action taken as a result of the events with diphenhydramine hydrochloride was post-therapy. Therapeutic measures were taken at the emergency room as a result of the events and included treatment with Topcid and Benadryl every 6 hours, 2x 40 mg of steroid. The information on the lot/batch number has been requested. Follow-up (02Jan2021): New information received from the same contactable healthcare professional reporting for herself includes: patient"s details, medical history, events updated, vaccine lot# and exp date, concomitant, seriousness of event "Her voice became raspy, she could hardly talk, she could barely talk" added as hospitalization, historical vaccine, suspect Benadryl details, therapeutic measures updated.; Sender"s Comments: A causal association between BNT162B2 and the reported events cannot be excluded based on a compatible temporal relation between vaccination and onset of events. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 934676 (history)
Form: Version 2.0
Age: 58.0

Sex: Female
Location: Washington
Vaccinated: 2020-12-31
Onset: 2021-01-02
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Arthralgia, Aspiration joint, Chills, Ultrasound joint
SMQs:, Arthritis (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Sertraline 25mg, Aspirin 81 mg, Calcium Citrate with Vitamin D3, Multivitamin with Iron, Hair Skin & Nails soft gel, Probiotic, Cetirizine
Current Illness: None
Preexisting Conditions:
Allergies: None
Diagnostic Lab Data: Ultrasound-guided aspiration of the right hip joint
CDC Split Type:

Write-up: Chills Hip pain

VAERS ID: 934749 (history)
Form: Version 2.0
Age: 38.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) /	- / 1	- / IM

PFIZER/BIONTECH		

Administered by: Unknown **Purchased by:** ?
Symptoms: Altered state of consciousness, Dizziness, Dyspnoea, Endotracheal intubation, Flushing, Hyperhidrosis, Immediate post-injection reaction, Intensive care, Mechanical ventilation, Pharyngeal swelling, Stridor, Tachycardia, Tachypnoea, Urticaria, Wheezing
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Asthma/bronchospasm (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypersensitivity (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Imitrex, Effexor XR, COVID-19 Vaccine (Pfizer)
Current Illness:
Preexisting Conditions: GERD, Migraines, Anxiety, allergic to Reglan
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: 38-year-old female who is healthcare worker and received first dose of COVID vaccine (Pfizer). Immediately after receiving the vaccine, patient developed lightheadedness, flushing, hives, wheezing and throat swelling. Patient was treated in an emergency department with epinephrine, gradually improved and was able to be sent home with an EpiPen, prednisone, hydroxyzine, and famotidine. The next day, patient again developed shortness of breath and her husband administered the EpiPen. EMS arrived and gave another dose of IM epinephrine and IV diphenhydramine. On arrival to the emergency department, the patient was altered, diaphoretic, tachypneic, tachycardic, and stridulous. Patient was given multiple doses of IM epinephrine and started on epinephrine drip. Stridor continued and was unresponsive to nebulized albuterol. Patient was then intubated and placed on mechanical ventilation. Other treatments included solumedrol, pepcid, magnesium sulfate, nebulized epinephrine, and IV fluids. admitted to the intensive care unit, weaned off epinephrine drip, and extubated the next day. Patient was monitored on hospital floor for one additional day and was then discharged with no residual symptoms.

VAERS ID: 934869 (history)

Form: Version 2.0
Age: 83.0
Sex: Female
Location: Michigan
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Body temperature increased, Condition aggravated, Facial paralysis, Hemiparesis, Rash erythematous, SARS-CoV-2 test negative, Skin warm
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow), Hearing impairment (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Prednisone 10mg, Donepezil HCL 10mg, Breo Ellipta inhaler, Spiriva Inhaler, Plavix
Current Illness: Rash, Non hemorrhagic CVA in 2009, Alzheimer''s dementia, COPD, HLD, CKD, Allergic Rhinitis
Preexisting Conditions: previous CVA, COPD, Alzheimer''s dementia
Allergies: none noted
Diagnostic Lab Data: Please see attached documentation
CDC Split Type:

Write-up: Patient had a rash prior to COVID vaccine. Prednisone 10mg started on 1/5/21 (only one dose administered) 1/6/21, 1am - Rash worsened with increase redness, warmth and extending of body surface, Temp 100.4, 1/6/21, 1:20am Benadryl 25mg administered, (Keflex was ordered at this time, however never administered). 1/6/21, 3am rash improved, temp 99.1 1/6/21, 6:50am Right facial droop and right sided weakness, sent to ER 1/6/21 transferred to hospital, continues to be hospitalized 1/11/2021

VAERS ID: 934889 (history)

Form: Version 2.0
Age: 93.0
Sex: Female
Location: Louisiana
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (UNKNOWN)) / UNKNOWN MANUFACTURER	- / UNK	- / -

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Arthralgia, Confusional state, Decreased appetite, Diarrhoea, Gait inability
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Pseudomembranous colitis (broad), Dystonia (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Arthritis (broad), Noninfectious diarrhoea (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: citalopram 20 mg PO DAILY clonazepam 0.5 mg PO DAILY lisinopril 10 MG PO DAILY loratadine 10 mg PO DAILY magnesium oxide 400 MG PO BID memantine 5 MG PO BID metoprolol tartrate 25 MG PO BID pantoprazole [Protonix] 40 MG PO DAILY warfarin 4
Current Illness:
Preexisting Conditions: Afib Anxiety Arthritis CAD (coronary artery disease) Cervical vertebral fusion CVA (cerebral vascular accident) Frequent UTI Gastritis Glaucoma Heart attack Hyperlipidemia Hypertension Hypomagnesemia Insomnia Pancreatic mass Presence of inferior vena cava filter Psoriasis Pulmonary embolus Spondylolisthesis
Allergies: chlordiazepoxide [From Librium] ciprofloxacin [From Cipro] codeine diazepam meperidine morphine Nitrofuran Analogues ondansetron [From Zofran] oxycodone Penicillins pentazocine [From Talwin] propoxyphene [From Darvon] Sulfa (Sulfonamide Antibiotics)
Diagnostic Lab Data:
CDC Split Type:

Write-up: HPI narrative: Patient was fine until 2 days ago. Patient does have chronic dementia but got Covid vaccine 11:00 2 days ago and that night seem to be confused and not able to walk since then. Patient poor appetite since yesterday. Patient had diarrhea 3 x 2 days ago. No vomiting no fever no cough does not appear to be short of breath. Patient also with

left hip pain for few days with no injury.All history obtained from caretaker at bedside.

VAERS ID: 934901 (history)
Form: Version 2.0
Age: 92.0
Sex: Male
Location: Ohio
Vaccinated: 2021-01-08
Onset: 2021-01-09
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	026L20A / 1	LA / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Abdominal pain
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: APAP, amlodipine, metoprolol, ramipril, spironolactone, dutasteride, vitamin B12,
Current Illness:
Preexisting Conditions: SCC, CHF, HTN, BPH, CARCINOMA OF COLON
Allergies: NKDA
Diagnostic Lab Data:
CDC Split Type:

Write-up: SEVERE ABDOMINAL PAIN

VAERS ID: 934906 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-22

Onset: 2021-01-04
Days after vaccination: 13
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / -

Administered by: Private **Purchased by:** ?
Symptoms: Amnesia, Aphasia, Blood pressure increased, Computerised tomogram, Confusional state, Low density lipoprotein increased, Magnetic resonance imaging, SARS-CoV-2 test negative, Transient ischaemic attack, Vision blurred
SMQs:, Dyslipidaemia (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Ischaemic central nervous system vascular conditions (narrow), Dementia (broad), Embolic and thrombotic events, arterial (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Glaucoma (broad), Hypertension (narrow), Lens disorders (broad), Retinal disorders (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: SHINGRIX
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cardiovascular disease, unspecified (family history of cardiovascular(CV) disease at young age); Cholesterol blood increased (elevated cholesterol w/lifestyle changes); Herpes simplex (Herpes simplex on lips); Menopause (post menopause); Non-smoker
Allergies:
Diagnostic Lab Data: Test Name: cholesterol; Result Unstructured Data: Test Result:elevated cholesterol w/lifestyle changes; Test Name: blood pressure; Result Unstructured Data: Test Result:elevated; Test Name: CT; Result Unstructured Data: Test Result:unknown results; Test Name: LDL; Result Unstructured Data: Test Result:192; Test Name: MRI; Result Unstructured Data: Test Result:unknown results; Test Date: 20210104; Test Name: Nasal Swab; Test Result: Negative ; Test Name: weight; Result Unstructured Data: Test Result:157
CDC Split Type: USPFIZER INC2021005404

Write-up: experienced symptoms of TIA; Severe aphasia; blurred vision; confusion; short term memory loss; elevated blood pressure; This is a spontaneous report from a contactable Other-HCP(Patient). A 57-year-old female patient received first dose of BNT162B2(PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration at right arm, on 22Dec2020 08:15 at single dose for COVID-19 immunization. The patient was not pregnant.

Medical history included herpes simplex on lips, post menopause, elevated cholesterol w/lifestyle changes. Known allergies reported as no. Concomitant medication included varicella zoster vaccine rge (cho) (SHINGRIX) for immunization. On 04Jan2021 08:30, the patient experienced symptoms of transient ischaemic attack(TIA), severe aphasia, blurred vision, confusion, short term memory loss, elevated blood pressure. The patient admitted to (Hospital name) (still here, hospitalization days reported as 2). Symptoms resolved except very mild aphasia. The patient had very few risk factors for TIA but did have family history of cardiovascular(CV) disease at young age, low density lipoprotein(LDL) was 192. The patient did not have diabetes, HTN, or known heart disease. She did not have severe anxiety. She did not smoke or use any substances. She walked about five miles 4x a week. Weigh reported as 157. Events reported as serious due to hospitalization. The patient had no Covid prior vaccination. Covid(nasal swab) was tested post vaccination on 04Jan2021, Covid test result was Negative. The event resulted in emergency room/department or urgent care. Treatment received for the adverse event included clopidogrel bisulfate(PLAVIX), acetylsalicylic acid (ASPIRIN), statin; potassium, CT, MRI, "telemet". The outcome of the events was recovering. Information on the lot/batch number has been requested.; Sender"s Comments: The Company cannot completely exclude the possible causality between the reported symptoms of transient ischaemic attack(TIA), severe aphasia, blurred vision, confusion, short term memory loss, elevated blood pressure and the administration of BNT162B2, based on the reasonable temporal association. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RA, Agency, as appropriate.

VAERS ID: 934912 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: Florida
Vaccinated: 2020-12-24
Onset: 2021-01-04
Days after vaccination: 11
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9291 / 1	LA / -

Administered by: Military **Purchased by:** ?
Symptoms: Computerised tomogram, Echocardiogram, Pain, Pelvic venous thrombosis, Ultrasound Doppler
SMQs:, Embolic and thrombotic events, venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Date: 202101; Test Name: CT scans; Result Unstructured Data: Test Result:unknown result; Test Date: 202101; Test Name: echo; Result Unstructured Data: Test Result:unknown result; Comments: and echo of her heart to make sure there is nothing else; Test Date: 202101; Test Name: doppler; Result Unstructured Data: Test Result:unknown result; Comments: doppler of bilateral legs
CDC Split Type: USPFIZER INC2021005480

Write-up: DVT; have pain in same site where DVT is; This is a spontaneous report from a contactable consumer. A 28-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number EK9291), via an unspecified route of administration in left deltoid on 24Dec2020 10:00 at first single dose for COVID-19 immunization. Medical history was none. There were no concomitant medications. Caller was calling to report a possible adverse reaction to the Pfizer Covid-19 vaccine. The patient was currently at hospital, she was admitted for deep vein thrombosis (DVT) of left iliac vein, the patient had no past history as to why this would happen, that she is only 28 years old. Received the vaccine on 24Dec2020, the following day she did have pain in same site where DVT was. Took ibuprofen for the pain. The patient was admitted yesterday 04Jan2020 for the DVT, they were currently treating her with Lovenox injections and prescribing dose for discharge is Eliquis. CT scans and three shots of Lovenox for it, doing a doppler of bilateral legs and echocardiogram (echo) of her heart to make sure there is nothing else. The AEs require a visit to emergency room. The patient was asking if she can still get the 2nd dose based off the adverse event she experienced. Outcome of DVT was not recovered, of pain was unknown.

VAERS ID: 934922 (history)
Form: Version 2.0
Age: 39.0
Sex: Male
Location: Texas
Vaccinated: 2020-12-16
Onset: 2020-12-21
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / -

Administered by: Private **Purchased by:** ?
Symptoms: Influenza like illness, Muscular weakness, Neurological symptom, Paraesthesia,

Peroneal nerve palsy, SARS-CoV-2 test negative
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Diabetes; Eosinophilic granulomatosis with polyangiitis; Hypertension; Migraine
Allergies:
Diagnostic Lab Data: Test Date: 20201219; Test Name: Nasal Swab; Test Result: Negative
CDC Split Type: USPFIZER INC2021006226

Write-up: Followed by neurological symptoms staring day 4; parasthesias of both upper extremity; progression to muscle weakness of all four extremities/Currently with significant muscle weakness, Left hand weakness leading to dropping of objects and left foot drop; progression to muscle weakness of all four extremities/Currently with significant muscle weakness, Left hand weakness leading to dropping of objects and left foot drop; Flu like symptoms first 3 days; This is a spontaneous report from a contactable physician (patient). A 39-year-old male patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot number: ek5730) at left arm, via an unspecified route of administration on 16Dec2020 at single dose for covid-19 immunisation. Medical history included hypertension, diabetes, migraines, Eosinophilic granulomatosis with polyangiitis (EGPA) remission. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient did not have any allergies to medications, food, or other products. The patient''s concomitant medications were not reported. On 21Dec2020, the patient experienced flu like symptoms first 3 days. Followed by neurological symptoms staring day 4, parasthesias of both upper extremity with progression to muscle weakness of all four extremities. Leading to 2 ER visits and hospital admission. Evaluation by internal medicine, neurology and rheumatology. Currently with significant muscle weakness, Left hand weakness leading to dropping of objects and left foot drop. The events resulted in Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Hospitalization, Disability or permanent damage. The treatment for events included High dose steroid. Covid test included Nasal Swab: negative on 19Dec2020. The outcome of events was not recovered.; Sender''s Comments: Based on temporal association, the causal relationship between bnt162b2 and the events influenza like illness, neurological symptom, paraesthesia, muscular weakness and peroneal nerve palsy cannot be excluded. The information available in this report is limited. This case will be reassessed once additional information becomes available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 934951 (history)
Form: Version 2.0
Age: 35.0
Sex: Male
Location: Arizona
Vaccinated: 2020-12-17
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / OT

Administered by: Private **Purchased by:** ?
Symptoms: Acute disseminated encephalomyelitis, Cerebrovascular accident, Diagnostic procedure, Dysarthria, HIV test, X-ray
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (narrow), Noninfectious encephalopathy/delirium (broad), Demyelination (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Hypertension
Allergies:
Diagnostic Lab Data: Test Name: HIV test; Result Unstructured Data: Test Result:Unknown results; Test Name: radiologic; Result Unstructured Data: Test Result:Acute demyelinating encephalomyelitis; Comments: states it looks like a radiologic diagnosis; Test Name: X-ray; Result Unstructured Data: Test Result:Unknown results
CDC Split Type: USPFIZER INC2021008606

Write-up: Acute demyelinating encephalomyelitis; Slurring his speech; Stroke; This is a spontaneous report from a contactable physician. A 35-year-old male patient received first dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), intramuscular in deltoid (unknown if right or left) on 17Dec2020 at 30 ug, single for "Preventative". Medical history included hypertension. There were no concomitant medications.(Physician) He is calling about the Pfizer Covid 19 vaccine. States what is going on with the patient may be associated

as a side effect. The patient got the vaccine two to three weeks ago, he clarifies the patient received the vaccine on 17Dec2020 and the patient ended up acutely developing (states it is a presumptive diagnosis) Acute demyelinating encephalomyelitis, states it looks like a radiologic diagnosis. The patient is an employee at hospital. When querying seriousness states it is medically significant but could be disabling but he thinks the patient will recover. Reporter seriousness for acute demyelinating encephalomyelitis: Medically significant, Hospitalization. Patient was hospitalized on Sunday and he is still admitted at this time. Dates when patient was in hospital for acute demyelinating encephalomyelitis was from 03Jan2021 to ongoing. Caller thinks the patient was flown to (Place) yesterday. The patient''s mother asked the caller if the caller thought the acute demyelinating encephalomyelitis was from the vaccine and the caller responded that he did not think it was from the vaccine. He confirms the patient is still admitted in the hospital and the patient''s attending neurologist is doctor. The caller heard about the patient from doctor. When querying covid vaccine dose, the caller states the standard dose is 30 mcg. This was clarified and documented as provided. The patient has not received his second dose yet. He asks if the patient should receive the second dose. He asks a general question if a pregnant patient can be given the Pfizer covid vaccine. He heard the patient had a stroke then the CFO tried to talk to him and the patient was slurring his speech. Caller spoke to the patient''s mother this morning and caller told the mother that he would try to find out what is going on with the patient. He asked that the patient get an HIV test even though he does not think the patient is at risk. Vaccination facility type was Hospital. Vaccine administered at military facility was No. None additional vaccines administered on same date of the PFIZER suspect. AE acute demyelinating encephalomyelitis require a visit to Emergency Room, not visit to physician office. Prior Vaccinations (within 4 weeks) was none. He has heard of acute demyelinating encephalomyelitis being associated with vaccines in the past and states that it is rare and usually in kids. States he saw patients that may have had acute demyelinating encephalomyelitis back in the 80s and 90s. Therapeutic measures were taken as a result of acute demyelinating encephalomyelitis (Patient will get steroids tonight pending the review of the x-ray). The outcome of the events was unknown. Information on the lot/batch number has been requested.; Sender''s Comments: The reported stroke with speech slurred, and the presumptive diagnosis of acute demyelinating encephalomyelitis (looks like a radiologic diagnosis by the reporting physician), was most likely an intercurrent disease, and unlikely causally related to the first dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE). The case will be reassessed should additional information become available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 934961 (history)
Form: Version 2.0
Age: 35.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-04
Onset: 2021-01-04
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Arthralgia, Blood glucose normal, Blood lactic acid decreased, Bone pain, Burning sensation, C-reactive protein increased, Chills, Crying, Fatigue, Grunting, Headache, Influenza virus test negative, Injection site erythema, Injection site pain, Injection site pruritus, Injection site swelling, Lymphadenopathy, Pain, Painful respiration, Pyrexia, SARS-CoV-2 test negative, Tachycardia, Tension headache, Vaccination complication
SMQs:, Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Depression (excl suicide and self injury) (broad), Osteonecrosis (broad), Arthritis (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Diabetes (Verbatim: Diabetes)
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20210105; Test Name: lactate; Result Unstructured Data: Test Result:1.5; Test Date: 20210105; Test Name: blood pressure; Result Unstructured Data: Test Result:90/50; Test Date: 202101; Test Name: blood pressure; Result Unstructured Data: Test Result:Normal range; Test Date: 20210104; Test Name: Temperature; Result Unstructured Data: Test Result:104; Test Date: 20210105; Test Name: Temperature; Result Unstructured Data: Test Result:102.8; Test Date: 202101; Test Name: Temperature; Result Unstructured Data: Test Result:105; Comments: axillary temperature; Test Date: 20210105; Test Name: CRP; Result Unstructured Data: Test Result:30; Test Date: 20210105; Test Name: heart rate; Result Unstructured Data: Test Result:144-152 at rest; Test Date: 202101; Test Name: heart rate; Result Unstructured Data: Test Result:109; Comments: normal range; Test Date: 20210105; Test Name: O2 sat; Result Unstructured Data: Test Result:95-96; Test Date: 20210105; Test Name: COVID-19 test; Test Result: Negative
CDC Split Type: USPFIZER INC2021010190

Write-up: Swollen lymph nodes; blood pressure was 90/50; Chills; body aches; Fatigue; Joint pain; Bone pain; Headache; Swelling, pain, redness and soreness at the injection site; Swelling, pain, redness and soreness at the injection site; Swelling, pain, redness and soreness at the injection site; Tachycardia / heart rate was 144-152; painful to breath and she was grunting; also said that the injection site itched a little bit.; Fever; First dose on 17Dec2020, second does on 04Jan2021; This is a spontaneous report from a contactable nurse reporting for herself. A 35-year-old female patient received the first single dose of

BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) ((lot number was unknown) intramuscularly on 17Dec2020 at 0.3ml, and the second single dose (lot number EJ1685) intramuscularly on 04Jan2021 07:45 at 0.3ml, for COVID-19 prophylaxis. Medical history included ongoing diabetes. Her diabetes was diagnosed about 6 months ago (reported on 06Jan2021) and was well controlled. Her glucose was at 140 at the ER (emergency room) even after drinking a drink that had sugar in it. The patient''s concomitant medications were not reported. The patient received the second dose on 04Jan2021 at 07:45 and experienced a severe reaction hours after she received it, and she ended up having to go the ER and that they were baffled about what happened. She reported her symptoms were: fatigue, joint pain mainly in her knees, bone pain that did not feel like muscle pain, headache, swollen lymph nodes in the left axilla, and swelling, redness, pain, and soreness in the left arm at the injection site and also said that the injection site itched a little bit. Her symptoms started about 9-10 at night on 04Jan2021. She said that she felt like with the second dose she noticed the soreness and pain and redness at injection site after about 2 hours after receiving the second dose, which was sooner than with the first injection. The fever started with the chills and body aches. The chills and body aches started at 22:00 on 04Jan2021. She took acetaminophen and melatonin to try and sleep it off. Fever was after that at around midnight 04Jan2021 and she was burning in fever and her temperature was 104. She stated she started taking layers off and got out from under her covers. She had a sweater on when she took her temperature and her axillary temperature was 105. She said that she got into a hot shower because that was the only thing that provided comfort to her chills. She had not had any chills in about the last 12 hours though (as reported on 06Jan2021). She said that the body aches were so severe that she just sat in her bed and cried. She had tachycardia and she was grunting in pain. She was rotating ibuprofen and acetaminophen and then her fever was not as high. She stated that if her fever came back it was lower each time, and if she did not take the medications though, the bone pain was excruciating. She said that with the bone pain it was like no-one can hold her hand or hug her. Fatigue started at around midnight 04Jan2021. Her headache was at midnight. She felt like she had a headband on and was radiating down the back of her neck. She noticed the lymph nodes were swollen after she went to the ER at around 07:00-08:00 05Jan2021. She was walking with pillow under her arm, and if she moved or lifted her arm it hurt. She went to the ER at 01:00 05Jan2021. Her blood pressure was 90/50, her heart rate was 144-152 at rest, temperature was 102.8, O2 saturation was 95-96. She said that it was painful to breath and she was grunting, but did not have any breathing issues. The nurse thought she was septic and notified the doctor. The nurse and the doctor did not think it was the vaccine and thought that she was COVID-19 positive. But after they tested her, she was negative for flu and COVID and they were baffled. They were unclear on what happened and did not think it was from the vaccine. They bolused her 1 liter of Normal Saline and gave her a dose of Fentanyl that minimally helped with the bone pain. She was also given 30mg of ketorolac (TORADOL) IV (intravenous) and that significantly improved her pain. She was there a total of 3 hours. She was admitted to the back area of the ER. She said that they drew labs and everything was within normal limits. Her CRP (C-reactive protein) was 30 and her lactate was at 1.5. She said that they told her that she was most likely having an immune response to the vaccine. Her heart rate and blood pressure came to a more normal range and everything returned to baseline. Her heart rate was 109. She was told to rotate paracetamol (TYLENOL) and ibuprofen for at least the next 24 hours and hydrate. The events fever, chills, fatigue, joint pain, bone pain, headache, swollen lymph nodes, and swelling, pain, redness and soreness at the injection site were serious due to hospitalization from 05Jan2021 to 05Jan2021. The patient was recovering from fever, chills, joint pain, headache, heart rate and blood pressure, not recovered from fatigue, bone pain, swollen lymph nodes and swelling, pain, redness and soreness at the injection site. The event swollen lymph nodes worsened. The outcome of "injection site itched a little bit" was unknown. The reporter considered the events fever, chills, fatigue, joint pain, bone pain, headache, swollen lymph nodes, and swelling, pain, redness and soreness at the injection site were all related to the vaccine; Sender''s Comments: Based on temporal association, the causal relationship between

BNT162B2 and the events pyrexia, chills, fatigue, pain, arthralgia, bone pain, headache, lymphadenopathy, vaccination site erythema, vaccination site swelling, vaccination site pain, blood pressure decreased, tachycardia, grunting and vaccination site pruritus cannot be excluded. The information available in this report is limited and does not allow a medically meaningful assessment. This case will be reassessed once additional information becomes available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 934990 (history)
Form: Version 2.0
Age: 26.0
Sex: Female
Location: Ohio
Vaccinated: 2021-01-07
Onset: 2021-01-09
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	012L20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Contusion, Immunoglobulin therapy, Injection site pain, Platelet count decreased, Platelet transfusion
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Accidents and injuries (narrow), Extravasation events (injections, infusions and implants) (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Ibuprofen and acetaminophen
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: 1/9/2021 18:30 2x10^3/uL 1/9/2021 21:30 6x10^3/uL 1/10/2021 02:10 8x10^3/uL 1/10/2021 07:30 18x10^3/uL 1/10/2021 18:12 127x10^3/uL 1/11/2021 05:43 142x10^3/uL

CDC Split Type:

Write-up: 26-year-old lady came in after she noticed she had bruises on her left hand after a CPR procedure at hospital. Patient was apparently in well health, she had received COVID-19 mRNA vaccine on January 7 at 3 PM, she has taken 2 pills with ibuprofen and tylenol for pain in right deltoid following vaccination. She was doing the CPR at 1:00 this afternoon, and she noticed that her left dorsum had some bruises. She took day off and went home and noticed that she also had bruises in both medial thighs, above the knee and some bruises in scalp. Patient presented to the Emergency Room 1/9/2021 ~6PM and platelet count was found to be 2x10^3/uL. Patient required transfusion of 7 units of platelets, steroids, and IVIG.

VAERS ID: 935266 (history)
Form: Version 2.0
Age: 41.0
Sex: Female
Location: Wisconsin
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal discomfort, Abdominal distension, Abdominal pain, Bowel movement irregularity, Nausea, Vomiting
SMQs:, Acute pancreatitis (broad), Retroperitoneal fibrosis (broad), Gastrointestinal perforation, ulcer, haemorrhage, obstruction non-specific findings/procedures (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient presents with abdominal pain that started in the middle of the night. Had first COVID vaccine the previous day. Patient states the pain is intermittent "comes and goes" "cramping" "pressure and bloating" feeling. Patient states her normal bowel movements are 12 times per day. The last time she went was this morning. She is concerned about an "obstruction" Patient states she has "some nausea" She states she has ate and drank normally today. Patient has a history of ulcerative colitis and a complete colectomy with a ileal rectal pouch. She has had abdominal pain since this morning which is crampy, associated with nausea and recurrent vomiting. She normally has 6-12 bowel movements a day, but none since this morning. She does feel her abdomen is distended as well. The last time she had anything like this was when she developed pouch itis last spring, but that was much less painful than this. Her appendix is gone, but she believes she still has her gallbladder.

VAERS ID: 935310 (history)
Form: Version 2.0
Age: 52.0
Sex: Female
Location: New York
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Paraesthesia
SMQs:, Peripheral neuropathy (broad), Guillain-Barre syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Went into the ED for bilateral hand and feet tingling. Worked up for possible

Guillain Barre.

VAERS ID: 935338 (history)
Form: Version 2.0
Age: 44.0
Sex: Male
Location: Kansas
Vaccinated: 2021-01-06
Onset: 2021-01-09
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Blood pressure increased, Blood test, Computerised tomogram head, Dizziness, Facial paralysis, Facial paresis, Feeling abnormal, Hypoaesthesia, Motor dysfunction, Muscular weakness, Paraesthesia, Paraesthesia oral, Speech disorder
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Akathisia (broad), Dyskinesia (broad), Dystonia (broad), Parkinson-like events (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Psychosis and psychotic disorders (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypertension (narrow), Hearing impairment (broad), Vestibular disorders (broad), Immune-mediated/autoimmune disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Hypertension, obesity, DM type II
Allergies: NKA
Diagnostic Lab Data: blood serum labs, CT brain on 1/9/21 at Hospital ER; scheduled for appt with PCP at on 1/12/21
CDC Split Type:

Write-up: Patient reported stroke-like symptoms as he was driving to work: started "feeling funny" with dizziness; progressed to feeling weakness in left side of face with facial drooping; tingling in left hand progressed to numbness and weakness in left arm and leg; difficultly coordinating motor movement in right arm; difficulty/ diminished speech "tongue felt fat."

Symptoms resolved within appx 1+ hour from onset. Per patient, he was given an injection at the hospital (unsure for what?) and was discharged with a prescription for chlorthalidone 25mg daily, blood pressure was elevated.

VAERS ID: 935350 (history)
Form: Version 2.0
Age: 81.0
Sex: Male
Location: Texas
Vaccinated: 2020-12-31
Onset: 2021-01-02
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Unresponsive to stimuli
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-06
Days after onset: 4
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Symbicort 160/4.5 mcg/act inhaler Dilitiazem ER 360mg PO daily Eliquis 5mg PO twice a day Furosemide 20mg PO daily Losartan 100mg PO daily KCL 10mEq ER PO daily Ranitidine 150mg PO Ventolin HFA 0.09mg/1ACT 2 puffs every 4 hours as needed Me
Current Illness: Dyspnea, Shortness of breath
Preexisting Conditions: Paroxysmal atrial fibrillation Hypertension Chronic iron deficiency anemia Congestive heart failure with chronic diastolic
Allergies: No known allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient was found unresponsive at home with SpO2 20% 1/2/2021

VAERS ID: 935452 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: New Mexico
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3248 / 2	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Diarrhoea, Echocardiogram normal, Electrocardiogram T wave abnormal, Myocarditis, Nausea, Platelet count decreased, Pyrexia, Thrombocytopenia, Troponin increased, Vomiting
SMQs:, Acute pancreatitis (broad), Haematopoietic thrombocytopenia (narrow), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Other ischaemic heart disease (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (narrow), Hypokalaemia (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Multivitamin daily
Current Illness: no
Preexisting Conditions: no
Allergies: NKDA
Diagnostic Lab Data: Troponin 0.08 and then 2.3 and up to 4 Platelets 85 and then 61.
CDC Split Type:

Write-up: 1/6/21 8pm started with Nasuea, vomiting, diarrhea and fever. 1/7/21 started having intermittent chest pain in the morning. Then in the evening it became constant. Went to ER that evening due to chest pain. EKG showed t wave abnormality. 1st Trop was negative went from 0.08 to 2.3 Had 2 Echo"s done and they were normal. Platelets were 85. Was discharged without chest pain. Troponin on discharge was 0.67 and platelets 61. Was admitted due to Chest pain and troponin. Attending provider diagnosed as myocarditis and

thrombocytopenia R/T vaccine.

VAERS ID: 935569 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Kansas
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 2	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal discomfort, Asthenia, Blood glucose increased, Chest pain, Diabetic ketoacidosis, Diarrhoea, Headache, Migraine, Pain, Pain in extremity, Pyrexia
SMQs:, Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Gastrointestinal perforation, ulcer, haemorrhage, obstruction non-specific findings/procedures (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Noninfectious diarrhoea (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Type 2 diabetes
Allergies: Sulfa: lip swelling Daptomycin: anaphylaxis Zosyn: lip swelling and hives Benadryl: Itching
Diagnostic Lab Data:
CDC Split Type:

Write-up: Pfizer COVID-19 Vaccine A couple hours after the vaccination the patient experienced pain in the vaccine arm, headache, and feeling ache. Day 1 post vaccination patient experienced sore arm, headache, low grade fever, feeling ache, and GI symptoms with diarrhea. Day 2 post vaccination patient experienced sore arm, Migraine, and diarrhea.

Day 3 post vaccine patient woke up with chest pain that radiated into her left arm and some weakness. Patient"s blood sugar was $g500 and was admitted to hospital for DKA.

VAERS ID: 935780 (history)
Form: Version 2.0
Age: 61.0
Sex: Female
Location: Ohio
Vaccinated: 2021-01-05
Onset: 2021-01-05
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain upper, Blood test, Chest X-ray, Computerised tomogram, Contusion, Dysstasia, Electrocardiogram, Eye contusion, Fatigue, Feeling of body temperature change, Headache, Loss of consciousness, Malaise, Memory impairment, Poor quality sleep, Sensation of foreign body, Swelling face, Swelling of eyelid, Tremor, Vaccination complication, Vision blurred
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dementia (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Accidents and injuries (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Glaucoma (broad), Lens disorders (broad), Retinal disorders (broad), Depression (excl suicide and self injury) (broad), Periorbital and eyelid disorders (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Levothyroxyn, Lisinopril
Current Illness: no
Preexisting Conditions: Thyroid
Allergies: Penicillin, Amoxicillin, Sulfur, pineapple

Diagnostic Lab Data: Ct Scan, EKG, chest xray, bloodwork
CDC Split Type:

Write-up: Extremely tired went to bed at 8:30. At 11:00, tossed and turned until 12:16 a.m. Did not think she could stand up. Walked in the bathroom had pain in stomach, was very hot and freezing at the same time, was shaking. Head hurt was hurting so bad. Tried standing up from toilet, was passed out. Eye, half of face is bruised and swollen , eye shut for two days. More than forty minutes passed before she could call her husband. Does not remember anything because she was passed out. Husband came in and grabbed her, to get her back into bed. she could not stand. Went to ER at Hospital. Had CT scan, blood work , EKG, chest xray. Was still kind of out of it but was able to communicate with nurses. Face is still black and blue. Last night got up at 4:00 a.m. something was wrong with her throat. Felt like a walnut was in her throat. Went to doctor this morning. They said it was a reaction from the shot. Got an EpiPen, prednisone. Call them in 2 days to let them know how she"s doing and also wear a heart monitor because they do not know how to take care of her, Has had vision blurriness in both eyes. Has not felt well at all.

VAERS ID: 935849 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-01
Onset: 2021-01-05
Days after vaccination: 35
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Appendicectomy, Appendicitis, Computerised tomogram abdomen, Computerised tomogram pelvis, Full blood count, Lymphadenopathy, Ultrasound pelvis
SMQs:, Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: None
Diagnostic Lab Data: CBC, pelvic ultrasound, CT abdomen and pelvis all on January 7,2021

CDC Split Type:

Write-up: Noted left lymph nodes left neck. On January 7, 10 days post injection had acute appendicitis requiring emergency appendectomy.

VAERS ID: 935984 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Colorado
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1685 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal discomfort, Blood test, Fatigue, Headache, Pyrexia, Urticaria
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Gastrointestinal perforation, ulcer, haemorrhage, obstruction non-specific findings/procedures (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Birth control
Current Illness: no
Preexisting Conditions: Lyme Disease, Heart Condition
Allergies: peanuts,
Diagnostic Lab Data: Given Benadryl, Beta Blocker, Steroid, blood work
CDC Split Type:

Write-up: About 15 minutes after vaccine, hr 155, fever 102, covered in hives, sick to her stomach and a pounding headache. Has had headache since then and been extremely fatigue.

VAERS ID: 936151 (history)
Form: Version 2.0
Age: 46.0
Sex: Female
Location: Montana
Vaccinated: 2020-12-16
Onset: 2020-12-23
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood test, Computerised tomogram head, Loss of proprioception, Magnetic resonance imaging, Magnetic resonance imaging brain, Magnetic resonance imaging spinal, Mobility decreased, Paraesthesia
SMQs:, Peripheral neuropathy (narrow), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Synthroid 6 am
Current Illness: R/O Covid infection, COVID test was negative December 4, 2020. Primary Care MD diagnosed me with COVID after symptoms consistent with the virus on December 14, 2020.
Preexisting Conditions: Hypothyroidism, Migraine headaches, IBS
Allergies: Morphine, extrapyramidal reaction to reglan
Diagnostic Lab Data: CT of brain X2, MRI of brain X2, MRI of C-Spine, MRI of brachioplexus, blood work.
CDC Split Type:

Write-up: One week after administration, I had sudden onset inability to move left arm. I was transported to ER immediately. Treated, scanned with CT of brain, MRI of brain, c-spine and brachioplexus. In hospital for 2 days and no answers. Still no answers to left arm paresthesia and proprioreceptor deficits. Spreading into left leg and mild systemic symptoms. I have been to the ER, seen by primary physician, Physiatrist and Neurology and Occupational Therapy. I am scheduled for many more appointments and trying to find and answer.

VAERS ID: 933233 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-20
Onset: 2020-12-20
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1688 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Angioedema, Anxiety, Dysgeusia, Medical observation, Paraesthesia oral, Respiratory distress, Tachycardia
SMQs:, Anaphylactic reaction (narrow), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Taste and smell disorders (narrow), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Acute central respiratory depression (broad), Guillain-Barre syndrome (broad), Hypersensitivity (narrow), Respiratory failure (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: INFLUENZA VIRUS;
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201220; Test Name: Medical observation; Result Unstructured Data: Test Result:unknown test results
CDC Split Type: GBPFIZER INC2021004559

Write-up: Angioedema; Respiratory distress; Taste metallic; Tingling mouth; Tachycardia; Anxiety; This is a spontaneous report from a contactable pharmacist downloaded from the Agency Regulatory Authority-WEB. This is a report received from the regulatory authority. The regulatory authority report number is GB-MHRA-WEBCOVID-20201220153356; Safety Report Unique Identifier: GB-MHRA-ADR 24545264. A 40-year-old female patient received the bnt162b2 (COMIRNATY; Lot Number: EJ1688), via an unspecified route of administration

on 20Dec2020 at a single dose for COVID-19 immunization. The patient medical history was not reported. The pharmacist reported the following: "the patient has not had symptoms associated with COVID-19. The patient has not been tested/or has had an inconclusive test for COVID-19. Unsure if patient is enrolled in clinical trial." Concomitant medications included influenza vaccine (MANUFACTURER UNKNOWN) taken for immunization, ramipril (MANUFACTURER UNKNOWN) taken for IgA nephropathy; both from an unspecified date to an unspecified date. The patient experienced the following events and outcomes: angioedema (hospitalization, medically significant, life threatening) on 20Dec2020 with outcome of unknown, respiratory distress (hospitalization, medically significant, life threatening) on 20Dec2020 with outcome of unknown, taste metallic (hospitalization, life threatening) on 20Dec2020 with outcome of not recovered, tingling mouth (hospitalization, life threatening) on 20Dec2020 with outcome of not recovered, tachycardia (hospitalization, life threatening) on 20Dec2020 with outcome of recovering, anxiety (hospitalization, life threatening) on 20Dec2020 with outcome of unknown. The patient stayed in the observation area for 15 minutes with no reaction. The patient was driving home and noted symptoms of metallic taste and tingling in mouth. The patient went back to the vaccination hub, and observations were taken at approximately 60 minutes after the injection. The patient was transferred to the local casualty for further management. The patient had no previous problems to vaccines. The patient already had a flu vaccine this year. The patient experienced the following symptoms: respiratory distress (20-30mins after vaccine), localised angioedema (20-30mins after vaccine), metallic taste in mouth, anxiety, tachycardia. The patient was treated with Intramuscular (IM) epinephrine (ADRENALINE) (2 doses), intravenous (IV) fluids, oxygen, chlorpheniramine/other antihistamines, hydrocortisone / other steroids. The events required hospitalisation. The patient had no previous allergic reaction. The patient underwent lab tests and procedures which included medical observation: unknown test results on 20Dec2020. Therapeutic measures were taken as a result of angioedema, respiratory distress, taste metallic, tingling mouth, tachycardia, and anxiety. No follow-up attempts are possible. No further information is expected.

VAERS ID: 934466 (history)
Form: Version 2.0
Age: 32.0
Sex: Male
Location: Foreign
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / 1	LA / -

Administered by: Other **Purchased by:** ?
Symptoms: Decreased appetite, Diarrhoea, Dizziness, Fatigue, Headache, SARS-CoV-2 test, Weight, Weight decreased
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad), Noninfectious diarrhoea

(narrow), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None
Allergies:
Diagnostic Lab Data: Test Date: 20210101; Test Name: COVID test nasal swab; Test Result: Negative ; Test Date: 20201230; Test Name: weight; Result Unstructured Data: Test Result:loss of 4 kg
CDC Split Type: ILPFIZER INC2021000752

Write-up: headaches; dizziness; fatigue; loss of appetite; diarrhea; lost around 4 kilos at that time; This is a spontaneous report from a contactable other hcp, the patient. This 32-year-old male patient received BNT162B2 (Pfizer-BioNTech COVID-19 mRNA vaccine), via an unspecified route of administration in the left arm on 30Dec2020 at 09:30 (at the age of 32-years-old) as a single dose for COVID-19 vaccination; and hpv vaccine vlp rl1 9v (yeast) (GARDASIL 9), via an unspecified route of administration in the left arm on 15Dec2020, dose number 3, as a single dose for immunization. The patient''s medical history was reported as none. The patient did not have any allergies to medications, food, or other products. Prior to the vaccination, the patient was not diagnosed with COVID-19. Concomitant medications were not reported. On 30Dec2020 at 09:45, the patient experienced having headaches, dizziness, fatigue, loss of appetite and diarrhea. He stated for 5 days, he couldn''t move from his bed, only sleeping and eating, that''s it. The patient had gone to the physician and the emergency room for the events. The patient reported he was in the ICU 3 times and they couldn''t find anything. He lost around 4 kilograms at that time. No therapeutic measures were taken as a result of the events. The clinical outcome of headaches, dizziness, fatigue, loss of appetite, diarrhea and weight loss was resolving. It was also reported that since the vaccination, the patient had been tested for COVID-19 on 01Jan2021, results were negative. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.; Sender''s Comments: Based on a compatible temporal relationship and currently known BNT162B2 vaccine safety profile causality between reported events and BNT162B2 vaccine cannot be completely excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

VAERS ID: 934762 (history)
Form: Version 2.0

Age:
Sex: Female
Location: Foreign
Vaccinated: 2021-01-02
Onset: 2021-01-02
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6797 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Anaphylactic reaction, Blood pressure abnormal, Blood pressure decreased, Blood pressure measurement, Chest discomfort, Dizziness, Dyspnoea, Flushing, Nausea, Pallor, Paraesthesia, Presyncope, Tachycardia, Tremor
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Parkinson-like events (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypertension (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (narrow), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Jessner"s lymphocytic infiltration
Allergies:
Diagnostic Lab Data: Test Date: 20210102; Test Name: Blood pressure; Result Unstructured Data: Test Result:80/40 mmHg
CDC Split Type: DKPFIZER INC2021013316

Write-up: Later lowblood pressure / Blod presure lowered to 80/40; Nausea; Flushing; tachycardia; Dizziness; shaking; affected blood pressure; Develop Difficulty breathing; Vasovagal reaction; Tingling in hands and feets; Anaphylactic reaction; Chest pressure; pale; This is a spontaneous report downloaded from the Agency Regulatory Authority-WEB, DK-DKMA-ADR 24551190. This is a spontaneous case, received on 02Jan2021 and follow up on 04Jan2021 from a physician and follow-up on 04Jan2021 from the hospital, which describes

the occurrence of Anaphylactic reaction (Anaphylactic reaction), Vasovagal attack, Blood pressure abnormal (affected blood pressure), Flushing (Flushing), Tachycardia (tachycardia) Blood pressure decreased (Later lowblood pressure / Blod presure lowered to 80/40), Nausea (nausea), Difficulty breathing (Develop Difficulty breathing), Chest tightness (chest pressure), Tingling feet/hands (Tingling in hands and feets), Shaking (shaking) and Pale (pale) in a 50-year-old female patient vaccinated with Comirnaty (tozinameran). There is no information regarding concomitant medication. There is no information regarding past medication. Patient concurrent conditions included Jessner''s lymphocytic infiltration. The patient have no former anaphylaxis. No CAVE noted. On 02Jan2021 the patient was vaccinated with first dose of Comirnaty 1 dosage form for COVID-19 immunisation. On 02Jan2021, 5 minutes later the patient developed Difficulty breathing On 02Jan2021, 10 minutes later the patient developed Nausea, Dizziness, Anaphylactic reaction, Blood pressure abnormal and Vasovagal attack. After 20 minutes the patient developed Chest tightness, Tingling feet/hands, Flushing, Blood pressure decreased and Tachycardia. On 02Jan2021 the patient developed Shaking and got Pale. The ADRs were by the physician reported as Life Threatening and resulting in hospitalisation(02Jan2021). The patient was treated with 0.3 mg adrenaline im., Tavegyl (clemastine) 2 mg x 1 and Solu-medrol (methylprednisolone) 80 mg followed by bolus sodium chloride 500 mL. The patient was send home with oral prednisolone for 3 days. Action taken with Comirnaty was unknown. The outcome of the events was recovered. The patient was observed for 6 hours at the emergency room. No progression of any symptoms. The patient was send home well-being. Test results: Blood pressure, 02Jan2021: 80/40 mmHg. FOLLOW-UP (version 002): Additional information received on 04Jan2021 from initial reporting physician: Comirnaty batchnumber is EJ6797 and solvens batch number 12PFL01. Route of administration was intramuscular. The ADRs were reportet as Life threatening. Reactions added: . The physician describes: Dizziness and nausea after 10 minutes. After further 10-15 minutes chest pressure and tingling in hands and feet. Gets blood pressure fall to 80/40. Calls 112. Immediate treatment for anaphylactic reaction with rp. Adrenaline 0.3 mg im. Green PVK in the left hand and gives tavegyl 2 mg x 1 and solu-medrol 80 mg followed by bolus nacl 500 mL. Upon arrival of the ambulance tachycardia and shaking as well as pale. Incidentally, stable vital parameters. FOLLOW-UP (version 003): Additional information received on 04Jan2021 from hospital. Woman familiar with morbus jessner, otherwise healthy. Develops difficulty breathing 5 minutes after administration of the vaccine and is admitted urgently to hospital. Getting better already in the emergency department, stable values. Admitted for observation for 6 hours and no progression of the symptoms Patient was send home well being with oral prednisolone for 3 days. Pt. has no CAVE.

VAERS ID: 934766 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-29
Onset: 2020-12-31
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6796 / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Asymptomatic COVID-19, Drug ineffective, Fracture, SARS-CoV-2 test
SMQs:, Lack of efficacy/effect (narrow), Accidents and injuries (narrow), Osteoporosis/osteopenia (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201231; Test Name: Covid 19 test; Test Result: Positive
CDC Split Type: EEPFIZER INC2021004609

Write-up: fracture of bone; Covid-19 infection, positive test result/asymptomatic; Covid-19 infection, positive test result/asymptomatic; This is a spontaneous report from a non-contactable physician. A 34-year-old female patient received the first dose bnt162b2 (COMIRNATY, lot number: EJ6796, expiry date: 21Apr2021), via an unspecified route of administration on 29Dec2020 at single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient experienced covid-19 infection, positive test result on 31Dec2020. The patient was diagnosed Covid-19 infection, positive test result came 31Dec2020. Patient was feeling well and was asymptomatic. Patient had positive Covid test after vaccine. They did test because, she turned to the emergency department for a fracture of bone. The outcome of events was unknown. The reporter didn''t think it was related with vaccination. No follow-up attempts are possible. No further information is expected. Follow-up (07Jan2021): New information received from a non-contactable physician included: new event (fracture of bone) and the reason for testing of the patient for covid-19 infection, patient''s information. No follow-up attempts are possible. No further information is expected.; Sender''s Comments: Based on the information available, a possible contributory role of the suspect BNT162B2 cannot be excluded for the reported asymptomatic Covid-19 infection based on the known safety profile. However the short duration of 2 days since the vaccine first dose is given it is unlikely patient would have fully developed immunity. The event fracture is most likely an intercurrent medical condition, due to an external factor, and is assessed as unrelated to BNT162B2.

VAERS ID: 934779 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-01

Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Dyspnoea, Heart rate, Malaise, Nausea, Palpitations
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ;
Current Illness: Atrial fibrillation (been well controlled with a heart rate of around 58-60 at baseline prior to taking the vaccine)
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Heart rate; Result Unstructured Data: Test Result:around 58-60 at baseline
CDC Split Type: GBPFIZER INC2021003946

Write-up: Palpitations; Malaise; Nausea; Shortness of breath; This is a spontaneous report from a contactable physician downloaded from THE regulatory authority-WEB. Regulatory authority report number GB-MHRA-EYC 00235986. Other case identifier number GB-MHRA-ADR 24545760. A 58-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, batch/lot number and expiration date unknown), intramuscularly on 15Dec2020 at single dose for COVID-19 immunisation. Medical history included had a background of ongoing atrial fibrillation but had been well controlled with a heart rate of around 58-60 at baseline prior to taking the vaccine. Concomitant medications included warfarin, simvastatin, dronedarone. The patient experienced palpitations on 16Dec2020; malaise, nausea and shortness of breath in Dec2020; all serious due to hospitalization. It was reported that after taking the COVID vaccine, the patient started developing palpitations on 16Dec2020, they were not too bad initially, but then 3 days later in Dec2020, the patient started feeling more unwell with shortness of breath, nausea and general malaise along with the palpitations and was admitted to the hospital at the time (unspecified date). The outcome of the event palpitations was not recovered, outcome of event shortness of breath was recovered on unspecified date, outcome of the other events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 934781 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-18
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Death, Pneumonia, Resuscitation, Sepsis
SMQs:, Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Sepsis (narrow), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2020-12-19
Days after onset: 1
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ;
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021009059

Write-up: Sepsis; Acute bronchopneumonia; This is a spontaneous report received from a contactable physician downloaded from the Regulatory Authority (GB-MHRA-EYC 00236063 and GB-MHRA-ADR 24546059). An 85-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), intramuscularly, on 15Dec2020 as a single dose for COVID-19 vaccination. The patient''s medical history was not reported. Concomitant medications included pregabalin (MANUFACTURER UNKNOWN), amitriptyline (MANUFACTURER UNKNOWN), amlodipine (MANUFACTURER UNKNOWN), candesartan (MANUFACTURER UNKNOWN), and levothyroxine (MANUFACTURER UNKNOWN). The patient experienced acute bronchopneumonia on 18Dec2020 and sepsis on an unspecified date. The events caused hospitalization and were reported as fatal. The clinical course was reported as follows: The patient was brought to the hospital by ambulance with severe sepsis and bronchopneumonia. She was resuscitated but unfortunately died shortly after arriving. The family reported that the patient received the coronavirus vaccine on 15Dec2020. It was

reported that it is unclear from the family history whether she was unwell before she received the vaccine. The clinical outcome of acute bronchopneumonia and sepsis was fatal. The patient died on 19Dec2020. The cause of death was reported as acute bronchopneumonia and sepsis. It was not reported if an autopsy was performed. No follow-up attempts are possible; information on batch number cannot be obtained.; Sender''s Comments: The information available in this report is limited and does not allow a medically meaningful assessment of the case. In particular the following relevant information is not available: medical history, autopsy report.; Reported Cause(s) of Death: Sepsis; Acute bronchopneumonia

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VAERS ID: 934783 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Lacunar stroke
SMQs:, Ischaemic central nervous system vascular conditions (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; CO-CODAMOL; ; ; ;
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021009604

Write-up: Taci stroke; This is a spontaneous report received from a contactable other healthcare professional by Pfizer from the Medicines and Healthcare products Regulatory Agency (UK-MHRA). Regulatory authority report number GB-MHRA-EYC 00236293, Safety report Unique Identifier GB-MHRA-ADR 24547108 and Sender''s case number ADR 24547108. A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH

COVID-19 VACCINE), via an unspecified route of administration on 22Dec2020 at single dose for COVID-19 immunisation. The patient medical history was not reported. Concomitant medications included bisoprolol, candesartan, clopidogrel, codeine phosphate, paracetamol (CO-CODAMOL), ibuprofen, lansoprazole, levothyroxine and simvastatin. The patient experienced total anterior circulation infarct stroke (TACI stroke) on an unspecified date, reported with seriousness criteria hospitalization. The outcome of the event was not recovered. No follow-up attempts are possible; information about batch/lot number cannot be obtained.

VAERS ID: 934786 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-19
Onset: 2020-12-27
Days after vaccination: 8
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Lacunar infarction
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; GANFORT; ; SENNA [SENNOSIDE A+B]
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021007148

Write-up: lacunar cerebral infarct; This is a spontaneous report from a contactable healthcare professional from the Regulatory Agency . The regulatory authority report number is GB-MHRA-EYC 00236937. A 92-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, on 19Dec2020

at a single dose for COVID-19 vaccination. The patient"s medical history was not reported. Concomitant medications included influenza vaccine (MANUFACTURER UNKNOWN) taken for influenza immunisation on 30Nov2020, carbomer (MANUFACTURER UNKNOWN), colecalciferol (MANUFACTURER UNKNOWN), ferrous fumarate (MANUFACTURER UNKNOWN), paracetamol (MANUFACTURER UNKNOWN), bimatoprost, timolol maleate (GANFORT), macrogol (MANUFACTURER UNKNOWN), and sennoside A+B (SENNA). The patient experienced lacunar cerebral infarct on 27Dec2020, which caused hospitalization and was reported as life-threatening. The clinical outcome of lacunar cerebral infarct was not recovered. No follow-up attempts are possible; information about batch/lot number cannot be obtained.; Sender"s Comments: The temporal relationship between the onset of the event and administration of the suspect product does not support a causal relationship.

VAERS ID: 934792 (history)
Form: Version 2.0
Age: 34.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-20
Onset: 2020-12-20
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJO724 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Oropharyngeal pain, Rash
SMQs:, Anaphylactic reaction (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021004349

Write-up: Rash; Throat pain; This is a spontaneous report downloaded from the Medicines Agency (MA) Regulatory authority-WEB GB-MHRA-WEBCOVID-20201220183940 and GB-MHRA-ADR 24545292. A contactable healthcare professional reported that a 34-year-old female patient received BNT162B2 (COMIRNATY, lot/batch ejo724), via an unspecified route of administration on 20Dec2020 at single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Patient has no medical history or known allergies to food, vaccines or medications. Patient has not had symptoms associated with COVID-19. Patient has not been tested/or has had an inconclusive test for COVID-19. Unsure if patient is enrolled in clinical trial. Patient is not pregnant. The patient experienced rash and throat pain (reported as hospitalized) on 20Dec2020. The patient received intravenous (IV) hydrocortisone and chlorphenamine for this. The outcome of the events was recovering. No follow-up attempts are possible. No further information is expected.

VAERS ID: 934796 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-20
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	JO533 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Computerised tomogram, Generalised tonic-clonic seizure
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Generalised convulsive seizures following immunisation (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ADCAL D3
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Partial seizures (Possible. Also reported in Sep2020. No treatment.)
Allergies:

Diagnostic Lab Data: Test Name: Computerised tomogram; Result Unstructured Data: Test Result:unknown test results
CDC Split Type: GBPFIZER INC2021003925

Write-up: Seizure grand mal, three full seizures; This is a spontaneous report from a contactable consumer downloaded from the Medicines Agency (MA) Regulatory authority-WEB regulatory authority number GB-MHRA-WEBCOVID-20201221083910, Safety Report Unique Identifier GB-MHRA-ADR 24545366. A 92-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), lot number JO533, via an unspecified route of administration on 15Dec2020 at single dose for COVID-19 immunisation. Medical history included partial seizures from Jun2020 (Possible. Also reported in Sep2020. No treatment.) Concomitant medication included insulin, simvastatin, acetylsalicylic acid, colecalciferol, calcium carbonate (ADCAL-D3). The patient experienced seizure grand mal on 20Dec2020, three full seizures. Admitted to hospital. The patient underwent lab tests and procedures which included computerised tomogram: unknown test results on an unspecified date. Outcome of event was recovering. Patient had not had symptoms associated with COVID-19. Patient had not been tested/or had had an inconclusive test for COVID-19. Patient was not enrolled in clinical trial. Patient had not tested positive for COVID-19 since having the vaccine. No follow-up attempts possible. No further information expected.

VAERS ID: 934802 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood glucose, Blood test, Body temperature, Chest X-ray, Computerised tomogram head, Culture urine, Delirium, Off label use, Product use issue, Pyrexia, SARS-CoV-2 test
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (narrow), Noninfectious meningitis (broad), Medication errors (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cholesterol high; Chronic obstructive pulmonary disease; Dementia Alzheimer"s type; Dyspepsia; Hypertensive; Ischaemic stroke; Stroke (with residual right sided weakness.); Suspected COVID-19 (Unsure when symptoms stopped); Type 2 diabetes mellitus; Weakness left or right side
Allergies:
Diagnostic Lab Data: Test Name: blood glucose; Result Unstructured Data: Test Result:unknown result; Test Name: Blood test; Result Unstructured Data: Test Result:normal; Comments: normal; Test Name: Body temperature; Result Unstructured Data: Test Result:37.8 Centigrade; Comments: 37.8 degrees; Test Name: Chest X-ray; Result Unstructured Data: Test Result:normal, unremarkable; Comments: normal, unremarkable; Test Name: Head CT; Result Unstructured Data: Test Result:normal, no acute intracranial abnormality; Comments: normal, no acute intracranial abnormality; Test Name: Urine culture; Test Result: Negative ; Comments: negative; Test Date: 20200421; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:Yes - Positive COVID-19 test; Comments: Yes - Positive COVID-19 test; Test Name: SARS-CoV-2 test; Test Result: Negative ; Comments: negative
CDC Split Type: GBPFIZER INC2021003926

Write-up: Fever/febrile at 37.8 degrees; Acute delirium; patient also received beclometasone dipropionate when receving BNT162B2; patient also received beclometasone dipropionate when receving BNT162B2; This is a spontaneous report received from a contactable physician downloaded from the Agency Regulatory Authority-WEB via Regulatory Authority (RA) GB-MHRA-WEBCOVID-20201221113316. Safety Report Unique Identifier GB-MHRA-ADR 24545462. An 86-year-old male patient received BNT162B2 (Pfizer BioNTech COVID vaccine), via an unspecified route of administration on 17Dec2020 at a single dose for COVID-19 vaccination. Medical history included suspected COVID-19 from 20Apr2020 (unsure when symptoms stopped), stroke from 2006 to an unknown date with residual right sided weakness, chronic obstructive pulmonary disease (COPD), ischaemic stroke, cholesterol high, hypertensive, Type 2 diabetes mellitus, dyspepsia, Dementia Alzheimer"s type. Concomitant medication included beclometasone dipropionate for chronic obstructive pulmonary disease (COPD), clopidogrel for ischaemic stroke, ezetimibe for cholesterol high, losartan for hypertensive, metformin for Type 2 diabetes mellitus, lansoprazole for dyspepsia, donepezil for Dementia Alzheimer"s type. All concomitants were taken from unknown date to unknown date. On 18Dec2020, the patient experienced fever, acute delirium onset approximately 20 hours after receiving first vaccine dose. This gentleman is a current in-patient under the care of the Geriatric team. He was admitted on 18Dec2020 after receiving his first vaccine dose the previous afternoon with acute confusion. He was noted to be febrile at 37.8 degrees in accident and emergency (A&E). He had a computerised tomography (CT) scan of his head which showed normal, no acute intracranial abnormality. Blood tests including a bone profile were normal. He was not found to be constipated and urine culture was negative, chest X-ray was unremarkable. They had found no clear cause for this man"s delirium and are concerned it may have been precipitated by his vaccine dose. The patient underwent lab tests and procedures which included blood glucose: unknown result, body temperature: 37.8 degrees centigrade on unknown date, chest x-ray: normal, unremarkable on unknown date, urine culture: negative on unknown date, COVID-19 virus test: yes - positive covid-19 test on 21Apr2020, sars-cov-2 test: negative on unknown date. Patient has not tested positive for COVID-19 since having the vaccine. Patient is not enrolled in clinical trial. The outcome of event fever was recovered on 18Dec2020, outcome of event acute

delirium was recovering. No follow-up attempts possible. No further information expected. Lot/Batch number cannot be obtained.

VAERS ID: 934810 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-15
Onset: 2020-12-15
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Chest pain, SARS-CoV-2 test
SMQs:, Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Coronary heart disease; Hypertension; Hypothyroidism; Type 2 diabetes mellitus
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2021004083

Write-up: Acute chest pain; This is a spontaneous report from a contactable pharmacist. This is a report received from the Medicines Agency (MA) Regulatory Authority-WEB. Regulatory authority or other manufacturer number (GB-MHRA-WEBCOVID-20201221135941). Safety Report Unique Identifier (GB-MHRA-ADR 24545655). A 57-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) batch/lot# unknown, via an unspecified route of administration on 15Dec2020 at single dose for COVID-19 vaccination. Medical history included type 2 diabetes mellitus, hypertension, hypothyroidism and coronary heart disease all from an unknown date and unknown if ongoing. Concomitant medication

included amlodipine (AMLODIPINE) for Hypertension, empagliflozin (EMPAGLIFLOZIN) for Diabetes, levothyroxine (LEVOTHYROXINE) for Hypothyroidism, indapamide (INDAPAMIDE) for Hypertension, gliclazide (GLICLAZIDE) for Diabetes, atorvastatin (ATORVASTATIN) for Coronary heart disease, sitagliptin (SITAGLIPTIN) for Diabetes; all used from an unknown date. The patient experienced acute chest pain (hospitalization, medically significant) on 15Dec2020 with outcome of recovered on 15Dec2020. Clinical course: Central chest pain 5-10 minutes after receiving the vaccine. Nurse did mention that patient became anxious before receiving the vaccine. Patient had not tested positive for COVID-19 since having the vaccine. Patient was admitted for observation ECG (electrocardiogram). The patient underwent lab tests and procedures which included COVID-19 virus test: no - negative covid-19 test. Unsure if patient had had symptoms associated with COVID-19 Patient was not enrolled in clinical trial. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 934811 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-17
Onset: 2020-12-17
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	NOT KNOWN / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood creatine phosphokinase, Blood creatine phosphokinase increased, Blood electrolytes, Blood electrolytes increased, Blood sodium, Blood sodium decreased, Blood urea, Blood urea increased, Chest X-ray, Computerised tomogram head, Fall, Loss of consciousness, SARS-CoV-2 test
SMQs:, Torsade de pointes/QT prolongation (broad), Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Retroperitoneal fibrosis (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Accidents and injuries (narrow), Hyponatraemia/SIADH (narrow), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; ; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Atrial fibrillation; Cerebrovascular accident NOS; Hypertension; Prostatism; Venous ulceration
Allergies:
Diagnostic Lab Data: Test Name: creatine kinase; Result Unstructured Data: Test Result:raised; Test Name: electrolytes; Result Unstructured Data: Test Result:raised; Test Name: sodium; Result Unstructured Data: Test Result:low; Test Name: urea; Result Unstructured Data: Test Result:raised; Test Name: Chest X-Ray; Result Unstructured Data: Test Result:no collapse, consolidation/oedema etc; Test Name: CT head; Result Unstructured Data: Test Result:no acute pathology; Test Date: 20201219; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2021003930

Write-up: Fall; collapse; raised creatine kinase; raised urea and electrolytes; raised urea and electrolytes; low sodium; This is a spontaneous report a contactable physician downloaded from the Medicines Agency (MA) Regulatory authority-WEB. Regulatory authority GB-MHRA-WEBCOVID-20201221143629, Safety Report Unique Identifier GB-MHRA-ADR 24545666. An 88-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not provided), via an unspecified route of administration on 17Dec2020 at single dose for COVID-19 immunization. Medical history included venous leg ulceration, atrial fibrillation, Previous cerebrovascular accident, Hypertension, and Prostatism. Concomitant medications included amlodipine for Hypertension, apixaban for Chronic atrial fibrillation, atorvastatin, losartan, omeprazole, paracetamol, and tamsulosin. The patient has not had symptoms associated with COVID-19. The patient is not enrolled in clinical trial. The patient received the vaccine on 17Dec2020. The patient experienced fall on 17Dec2020. He was found collapsed at home by daughter on 17Dec2020. The next morning, 18Dec2020, the patient was admitted to the hospital. It was treated as fall with long lie. The physician stated that it was quite possibly unrelated to vaccine but given proximity of timing the physician felt it was necessary to report. The patient tested negative for COVID-19 on 19Dec2020 (reported as since having the vaccine). The patient underwent lab tests and procedures which included CT head - no acute pathology, chest x-ray - no collapse, consolidation/oedema etc, bloods show raised creatine kinase, raised urea and electrolytes and low sodium on an unspecified date. The outcome of the events was unknown. No follow-up attempts are possible, information on batch number cannot be obtained.; Sender''s Comments: The information currently provided is too limited to make a meaningful medical assessment hence, the events are conservatively assessed as related to the suspect drug BNT162B2 until further information becomes available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 934821 (history)
Form: Version 2.0
Age:

Sex: Female
Location: Foreign
Vaccinated: 2020-12-20
Onset: 2020-12-21
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident, Facial paresis, Hemiparesis, SARS-CoV-2 test, Vaccination site pain
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow), Immune-mediated/autoimmune disorders (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ;
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201202; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:Negative; Comments: No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2021003976

Write-up: Weakness left or right side; left side weakness (face); Vaccination site tenderness; Stroke; This is a spontaneous report from a contactable consumer downloaded from the Medicines Agency (MA) Regulatory authority-WEB GB-MHRA-WEBCOVID-20201221182228, Safety Report Unique Identifier GB-MHRA-ADR 24545822. An 88-year-old female patient received BNT162B2 (COMIRNATY), via an unspecified route of administration on 20Dec2020 at a single dose for COVID-19 immunization. The patient"s medical history was not reported. Concomitant medications included furosemide for Blood pressure high, gabapentin for Pain relief, and enalapril for Blood pressure high. The patient experienced stroke on 21Dec2020, weakness left or right side, left side weakness (face) on an unspecified date, and vaccination site tenderness on 21Dec2020. Patient has not had symptoms associated with COVID-19. Patient had vaccine on 20th December, had tenderness in area of vaccine, suffered suspected stroke on 21st December, left side weakness (face, arm), admitted to hospital.

Patient has not tested positive for COVID-19 since having the vaccine. Patient is not enrolled in clinical trial. The patient underwent lab tests and procedures which included COVID-19 virus test: No - negative COVID-19 test on 02Dec2020. The outcome of the event stroke was not recovered while the outcome of the other events was unknown. No follow-up attempts are possible; information about lot/batch number can not be obtained.

VAERS ID: 934829 (history)
Form: Version 2.0
Age: 51.0
Sex: Male
Location: Foreign
Vaccinated: 2020-12-21
Onset: 2020-12-21
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Allergic oedema, Eye pruritus, Eye swelling, Throat irritation
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Bronchiectasis; Drug intolerance (possible mushroom products); Comments: Unsure if patient has had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021009055

Write-up: Itchy eyes; Eye swelling; Itchy throat; Allergic oedema; This is a spontaneous report received from a contactable physician by Pfizer from the Regulatory Agency (RA). The regulatory authority report number is GB-MHRA-WEBCOVID-20201222123514, Safety

Report Unique Identifier GB-MHRA-ADR 24546166. A 51-year-old male patient received bnt162b2 (Pfizer-BioNTech COVID-19 mRNA vaccine), via an unspecified route of administration on 21Dec2020 at single dose for COVID-19 vaccination. Unsure if patient has had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial. Medical history included drug intolerance (possible mushroom products) and bronchiectasis. The patient"s concomitant medications were not reported. The patient previously took diltiazem and experienced drug intolerance. The patient experienced itchy eyes, eye swelling and itchy throat on an unspecified date and allergic oedema on 21Dec2020. The events were reported as medically significant. Swelling surrounding eyes, itchy eyes, scratchy throat. Not resolving with chlorphenamine. Presented to hospital. No sign of anaphylaxis. Reaction not worsening. Received intravenous (IV) hydrocortisone and chlorphenamine. Patient has not tested positive for COVID-19 since having the vaccine. The reporter saw this patient in the emergency department and they were discharged home at the time, so did not have additional information on Recovery since then. At the time the reporter saw them (the day after the vaccination) their symptoms had stabilised, not getting worse or better. From memory, the onset of symptoms was that they had itching of the eyes very quickly e.g. within 15-30 minutes of the vaccination, periorbital swelling a few hours later, and then a scratchy feeling of the throat during the night following it. The reporter did not diagnose anaphylaxis and they did not receive adrenaline. The diagnosis was "allergic reaction", and they were discharged with oral antihistamines. The patient had a possible reaction to mushrooms earlier in life so had been avoiding them since. Therapeutic measures were taken as a result of the events. The outcome of event allergic oedema was recovering. The outcome of other events was not recovered. No follow-up attempts are possible, information about lot/batch number cannot be obtained.

VAERS ID: 934831 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-18
Onset: 2020-12-19
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ0553 / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Circulatory collapse, Hypertension, Peripheral swelling, SARS-CoV-2 test
SMQs:, Cardiac failure (broad), Anaphylactic reaction (narrow), Angioedema (broad), Neuroleptic malignant syndrome (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Torsade de pointes, shock-associated conditions (narrow), Hypovolaemic shock conditions (narrow), Toxic-septic shock conditions (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypoglycaemic and neurogenic shock conditions (narrow), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypertension (narrow),

Hypersensitivity (narrow), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201221; Test Name: blood pressure; Result Unstructured Data: Test Result:high; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:not tested positive; Test Date: 20200909; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:Yes - Positive COVID-19 test
CDC Split Type: GBPFIZER INC2021008947

Write-up: Blood pressure high; Collapsed; Swollen arm; This is a spontaneous report from a contactable other health care professional downloaded from the Medicines Agency (MA) Regulatory Authority- WEB GB-MHRA-WEBCOVID-20201222135025, Safety Report Unique Identifier GB-MHRA-ADR 24546203. A 55-year-old female patient received BNT162B2 (COVID-19 MRNA VACCINE BIONTECH, batch/lot number EJ0553), via an unspecified route of administration on 18Dec2020 at single dose for COVID-19 vaccination. The patient''s medical history and concomitant medications were not reported. The patient experienced swollen arm on 19Dec2020 with outcome of not recovered, blood pressure high and collapsed both on 21Dec2020 with outcome of recovered on 21Dec2020. All events were reported as serious (hospitalization). The patient underwent lab tests and procedures which included COVID-19 virus: yes - positive covid-19 test on 09Sep2020. Patient had not had symptoms associated with COVID-19. Patient was not enrolled in clinical trial. Patient had not tested positive for COVID-19 since having the vaccine. No follow-up attempts possible. No further information expected.

VAERS ID: 934832 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-29
Onset: 2020-12-30
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	NOT KNOWN / 2	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Cellulitis, Erythema, Pyrexia, Rash, SARS-CoV-2 test, Sensitive skin, Swelling, Swelling face
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Test Result: Negative ; Comments: No - Negative COVID-19 test
CDC Split Type: GBPFIZER INC2021009508

Write-up: Cellulitis; felt like sunburn and was uncomfortable; Face red; Rash; Facial swelling; swelling started to spread into her neck; Slight temperature; This is a spontaneous report received from a contactable other health professional by Pfizer from the Regulatory Agency (RA). The regulatory authority report number is GB-MHRA-WEBCOVID-20201231115747. A 55-year-old female patient received second dose of bnt162b2 (COVID-19 MRNA VACCINE BIONTECH, batch/lot number not known, with unknown expiration date), via an unspecified route of administration on 29Dec2020 at single dose for covid-19 vaccination. The patient''s medical history and concomitant medications were not reported. The patient is well with no underlying conditions, or history of allergic reaction to vaccines. Patient has not had symptoms associated with COVID-19. Patient is not enrolled in clinical trial. The patient previous received first dose of bnt162b2 via an unspecified route of administration on 08Dec2020 at single dose for covid-19 immunization with no side effects. The patient experienced face red, facial swelling, slight temperature and rash, all on 30Dec2020, cellulitis on an unspecified date. The patient had her first vaccine on 08Dec2020 in the afternoon with no side effects. The second dose was given on the 29Dec2020 in the afternoon and on the evening of the 30Dec2020, a red rash appeared on her face. This got ''angrier'' and spread across her face, through the night, and swelling started to spread into her neck. She attended the urgent care center of the morning of the 31Dec2020. She stated that it was not painful, it felt like sunburn and was uncomfortable. The urgent care center noted a slightly raised temperature. Patient has not tested positive for COVID-19 since having the vaccine. This was the second dose of vaccine. The member of staff attended the urgent care center and received antibiotics and antihistamine tablets. Follow up received 04Jan2021: Patient admitted to hospital and doctors believe this to be facial cellulitis. The patient underwent lab tests and procedures which included COVID-19 virus test: no- negative on an unspecified date. The outcome of events for slight temperature was resolving, for cellulitis was unknown,

for other events was not resolved. No follow-up attempts are possible, information on batch number cannot be obtained.

VAERS ID: 934840 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-22
Onset: 2020-12-24
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Blood test, Dehydration, Diarrhoea, Dizziness, Oxygen saturation decreased, Physical examination, SARS-CoV-2 test, X-ray
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Anticholinergic syndrome (broad), Pseudomembranous colitis (broad), Acute central respiratory depression (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad), Noninfectious diarrhoea (narrow), Respiratory failure (broad), Infective pneumonia (broad), Dehydration (narrow), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Blood test; Result Unstructured Data: Test Result:unknown results; Test Name: Physical examination; Result Unstructured Data: Test Result:unknown results; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:No - Negative COVID-19 test; Test Name: X-ray; Result Unstructured Data: Test Result:unknown results
CDC Split Type: GBPFIZER INC2021010015

Write-up: Dizziness/lightheaded; severe diarrhoea; dehydration; Severe diarrhoea and low oxygen levels. Dehydration; This is a spontaneous report received from a contactable

consumer by Pfizer from the Medicines and Healthcare products Regulatory Agency (MHRA). The regulatory authority report number is GB-MHRA-ADR 24550290. An 87-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 22Dec2020 at single dose for COVID-19 vaccination. The patient medical history was not reported. Concomitant medication included influenza vaccine (INFLUENZA VIRUS) in Nov2020, omeprazole, STATINS. The patient experienced severe diarrhoea and low oxygen levels. dehydration (hospitalization, disability, medically significant) on 24Dec2020, dizziness (hospitalization, disability, medically significant) on an unspecified date. Patient had not had symptoms associated with COVID-19. Patient was not enrolled in clinical trial. Severe diarrhoea for nearly two weeks, lightheaded, dehydration, Low oxygen reading. Patient had not tested positive for COVID-19 since having the vaccine. The patient underwent lab tests and procedures which included the result of blood test was unknown; the result of physical examination was unknown; the result of COVID-19 virus test was no - negative covid-19 test; the result of x-ray was unknown. The outcome of the event dizziness/ lightheaded was unknown. The outcome of other event was not recovered. No follow-up attempts possible. No further information expected.

VAERS ID: 934855 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-29
Onset: 2020-12-31
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Balance disorder, Confusional state, Hallucination
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Psychosis and psychotic disorders (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Vestibular disorders (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: MADOPAR

Current Illness: Parkinson"s disease; Partial sight (partially sighted)
Preexisting Conditions: Comments: Patient has not had symptoms associated with COVID-19 Patient has not been tested/or has had an inconclusive test for COVID-19. Patient is not enrolled in clinical trial
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021009871

Write-up: Hallucinating; Falling; Confusional state; This is a spontaneous report from a contactable consumer. This is a report received from the Medicines and Healthcare products Regulatory Agency (MHRA). The regulatory authority report number is GB-MHRA-WEBCOVID-20210103111852. An 81-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 29Dec2020 as single dose for COVID-19 immunization and benserazide hydrochloride, levodopa (MADOPAR), via an unspecified route of administration from an unspecified date at 12 mg for Parkinson"s disease. Medical history included ongoing visual impairment, reported as partially sighted and ongoing Parkinson"s disease since unspecified dates. The patient"s concomitant medications were not reported. The patient experienced hallucinating, falling and confusional state on 31Dec2020, which were serious by the reporter as they were life threatening, involved hospitalization, caused disability, and was medically significant. Details were as follows: Patient has not had symptoms associated with COVID-19 and has not been tested, nor has he had an inconclusive test for COVID-19. Patient is not enrolled in a clinical trial. The patient was fine and coping with Parkinson"s and being partially sighted as best he could (per the reporter), but on the 29th he had his COVID vaccine and on the 31th, degenerated so quickly and badly that he is now in the hospital. He"s hallucinating, making no sense, has no idea where he is and lost all balance and has fallen more than twice. The outcome of hallucinating was not recovered; the outcomes of falling and confusional state were unknown. No follow-up attempts are possible; information about batch/lot number cannot be obtained.

VAERS ID: 934867 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2021-01-03
Onset: 2021-01-04
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident, Computerised tomogram, Hypertension
SMQs:, Neuroleptic malignant syndrome (broad), Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous

(narrow), Hypertension (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 202101; Test Name: CT examination; Result Unstructured Data: Test Result:results unknown
CDC Split Type: ILPFIZER INC2021009543

Write-up: blood pressure was high; The diagnosis was stroke / felt weakness in the hand and leg and dizziness; This is a spontaneous report from contactable consumer. A 66-year-old male patient received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 03Jan2021 at single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient was vaccinated on Sunday (03Jan2021). On Monday (04Jan2021) he felt weakness in the hand and leg and dizziness. The blood pressure was high and due to that, he was sent to hospital from the clinic in ambulance. He received anticoagulation treatment and underwent CT examination. Later, his situation got better but he was still in hospital for observation. The diagnosis was stroke. It was reported that there was no connection between the event to the vaccine and therefore, he was planning to take the second dose as well. The outcome of events was unknown. No follow-up attempts are possible, information about lot number cannot be obtained.

VAERS ID: 934875 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-27
Onset: 2020-12-27
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -

Administered by: Other **Purchased by:** ?

Symptoms: Off label use, Paresis, Product use issue, Pyrexia, Suppressed lactation
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad), Functional lactation disorders (narrow), Medication errors (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: PLPFIZER INC2021004726

Write-up: She reported reduced lactation; fever; hand paresis; Breastfeeding patient - vaccinated with the first dose on 27Dec2020.; Breastfeeding patient - vaccinated with the first dose on 27Dec2020.; This is a spontaneous report from a contactable pharmacist. This pharmacist reported similar events for 5 mothers. This is the fourth of 5 maternal cases. A female breastfeeding mother of an unspecified age received her first dose of bnt162b2 (COMIRNATY; lot number and expiry date was not reported) via an unspecified route of administration on 27Dec2020 at single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient experienced reduced lactation, fever and hand paresis on an unspecified date. It was also reported that the breastfeeding patient was vaccinated with the first dose of bnt162b2 on 27Dec2020. The patient was hospitalized due to the events reported on an unspecified date. Outcome of the events was unknown. Information about lot number and expiry date for the suspect product will be requested in follow-up attempts.; Sender''s Comments: Based on the reasonable temporal association and lacking alternative explanations, the Company cannot completely exclude the possible causality between the reported reduced lactation, fever and hand paresis and the administration of bnt162b2 for this female breastfeeding mother. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RA, IEC, as appropriate.,Linked Report(s) : PL-PFIZER INC-2021004718 same reporter/drug, similar events, different patients;PL-PFIZER INC-2021004729 child case

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VAERS ID: 936715 (history)
Form: Version 2.0

Age: 24.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2021-01-11
Onset: 2021-01-11
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Dyspnoea, Intensive care, Palpitations, Vomiting
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Vestibular disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Sprintec, QVar ProAir, Prevacid, Respi Click
Current Illness: Unknown
Preexisting Conditions: Asthma
Allergies: Sulfa, Neomycin, Bacitracin, Motrin, Shellfish, Cats/Dogs
Diagnostic Lab Data: Would need to obtain intensive care record from the hospital.
CDC Split Type:

Write-up: Approx 10-15 post vaccine, employee said she felt lightheaded and like her heart was racing. Within 10 minutes she said she felt difficulty breathing, She then vomited. The observation nurse at the clinic administered Epi Pen and called a Code. The employee was transported to the Emergency Dep"t and then to intensive care. She was placed on an Epi drip.

VAERS ID: 936891 (history)
Form: Version 2.0
Age: 64.0
Sex: Male
Location: North Dakota

Vaccinated: 2021-01-08
Onset: 2021-01-09
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL1284 / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Asthenia, Fall, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Accidents and injuries (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 4 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Increased weakness leading to a fall and fever of 101.3

VAERS ID: 936993 (history)
Form: Version 2.0
Age: 42.0
Sex: Female
Location: Iowa
Vaccinated: 2020-12-31
Onset: 2020-12-31
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	LA / IM

Administered by: Private **Purchased by:** ?

Symptoms: Acute myeloid leukaemia, Atypical pneumonia, Back pain, Biopsy bone marrow abnormal, Blast cells present, Blood smear test, C-reactive protein, Chills, Computerised tomogram thorax abnormal, Fatigue, Fibrin D dimer, Full blood count, Metabolic function test, Pain, Pyrexia, Red blood cell sedimentation rate, SARS-CoV-2 test negative, White blood cell count increased, White blood cell morphology abnormal
SMQs:, Haematopoietic cytopenias affecting more than one type of blood cell (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Blood premalignant disorders (narrow), Malignancy related therapeutic and diagnostic procedures (narrow), Cardiomyopathy (broad), Malignant lymphomas (broad), Myelodysplastic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Haematological malignant tumours (narrow), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 7 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Levothyroxine, Trintellix
Current Illness: Strep
Preexisting Conditions: Thyroid cancer, hypothyroidism, GERD
Allergies: Compazine
Diagnostic Lab Data: 1.5.2020: CBC, CMP, D Dimer, CRP, ESR. CTA of chest negative for PE but demonstrated atypical Pneumonia. RAPID covid test negative. Peripheral smear demonstrated blasts, auer rods. 1.6.2020: Bone marrow biopsy confirmed AML
CDC Split Type:

Write-up: Patient began experiencing fevers, body aches, back pain, fatigue, chills the night she received the vaccine. Symptoms progressed for the following four days when she was ultimately seen in PCP office. Laboratory evaluation demonstrated atypical pneumonia and elevated WBC count. Patient was diagnosed with Acute Myeloid Leukemia via bone marrow biopsy and is receiving treatment at the hospital.

VAERS ID:	937089 (history)
Form:	Version 2.0
Age:	50.0
Sex:	Female
Location:	Massachusetts
Vaccinated:	2021-01-07
Onset:	2021-01-08
Days after vaccination:	1
Submitted:	0000-00-00
Entered:	2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site /

		Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9321 / 2	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Balance disorder, Burning sensation, Head discomfort, Headache, Hypoaesthesia, Magnetic resonance imaging, Paraesthesia
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Vestibular disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Augmentin, Wellbutrin, Duloxetine, Klonipin
Current Illness: Diverticulitis 24 hrs prior
Preexisting Conditions: Fibromyalgia, mild asthma, anxiety and depression, occasional Diverticulitis, occasional headache/migraines
Allergies: n/a
Diagnostic Lab Data: MRI
CDC Split Type:

Write-up: Within 24 hrs, developed headaches, burning sensation down spine and neck. fullness in head intensified next day. balance was off, numbness and tingling throughout body. Went to ER. Diagnosed Paresthesia

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VAERS ID: 937343 (history)
Form: Version 2.0
Age: 37.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-07
Onset: 2021-01-07
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Abdominal pain, Dizziness, Endoscopic retrograde cholangiopancreatography, Headache, Hepatic enzyme increased, Nausea, Rash, Rash erythematous
SMQs:, Liver related investigations, signs and symptoms (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis

(broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Vestibular disorders (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: unknown
Diagnostic Lab Data: Per her supervisor, her "liver enzymes are elevated/out of whack" and she was having an ERCP
CDC Split Type:

Write-up: Within approximately 30 minutes after vaccine (Thursday), patient presented with red rash to upper chest, severe headache, dizziness and nausea. Was treated with Benadryl and Tylenol per anesthesia onsite. Symptoms remained throughout the day with minimal improvement. The following morning (Friday) only headache remained. patient began having abdominal pain, and was admitted to hospital the next day (Saturday). It is now Tuesday and patient is still hospitalized. Unclear if this entire event is vaccine related.

VAERS ID: 937474 (history)
Form: Version 2.0
Age: 28.0
Sex: Male
Location: Puerto Rico
Vaccinated: 2021-01-11
Onset: 2021-01-12
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Military **Purchased by:** ?
Symptoms: Chills, Hypoaesthesia, Malaise, Nausea, Pyrexia
SMQs:, Acute pancreatitis (broad), Peripheral neuropathy (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Sexual dysfunction (broad)

Life Threatening? No

Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: Dust, avocados
Diagnostic Lab Data: None.
CDC Split Type:

Write-up: Pt experienced the following morning: fever (100 F) chills, malaise, nausea, and numbness in hands at 0624 and 0836. Pt went to hospital at 1000, tx with IV fluids.

VAERS ID: 937480 (history)
Form: Version 2.0
Age: 85.0
Sex: Female
Location: Louisiana
Vaccinated: 2021-01-09
Onset: 2021-01-11
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Anxiety, Blood magnesium normal, Cardiovascular evaluation, Computerised tomogram head normal, Full blood count normal, Headache, Metabolic function test normal, Muscular weakness, Neck pain, Paraesthesia
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Noninfectious encephalopathy/delirium (broad), Arthritis (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes

Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Losartan,Amlodipine,Levothyroxine, Metoprolol, Crestor
Current Illness: None reported
Preexisting Conditions: Hypertension, hypothyroidism, hyperlipidemia
Allergies: None reported
Diagnostic Lab Data: As above on 1/11/2021
CDC Split Type:

Write-up: In the early morning of Monday, January 11th the patient developed a significant headache with neck pain. She also reported parathesia and tingling in bilateral upper extremities with weakness of the right upper extremity. She reported feeling very anxious and "wound up". Patient presented to the Emergency Department 3 hours later. CT of the head/brain, EKG, CBC,CMP, Magnesium and Cardiac profile were performed with no significant findings. Ativan 0.5mg was administered orally. Patient was admitted to the facility for observation. Symptoms gradually resolved with no additional treatment.

VAERS ID: 937579 (history)
Form: Version 2.0
Age: 64.0
Sex: Male
Location: Minnesota
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Abdominal pain, Acute respiratory distress syndrome, Arteriosclerosis coronary artery, Ascites, Atrial fibrillation, Blood culture negative, Blood lactic acid, Blood potassium decreased, Brain natriuretic peptide normal, C-reactive protein increased, Chronic obstructive pulmonary disease, Computerised tomogram abdomen abnormal, Computerised tomogram thorax abnormal, Confusional state, Culture urine positive, Diarrhoea, Dyspnoea, Electrocardiogram abnormal, Fall, Haematocrit decreased, Haemoglobin decreased, Hepatic cirrhosis, Hiatus hernia, International normalised ratio increased, Lipase normal, Lung opacity, Myalgia, Nausea, Oedema, Pancreatitis, Platelet count decreased, Portal hypertension, Procalcitonin increased, Pulmonary embolism, Pyrexia, Red blood cell count decreased, Scan with contrast, Sepsis, Sinus tachycardia, Splenic granuloma, Tachycardia, Tachypnoea, Vomiting, White blood cell count decreased
SMQs:, Rhabdomyolysis/myopathy (broad), Cardiac failure (broad), Liver related investigations, signs and symptoms (narrow), Hepatic failure, fibrosis and cirrhosis and other liver damage-related conditions (narrow), Liver-related coagulation and bleeding disturbances (narrow), Anaphylactic reaction (narrow), Acute pancreatitis (narrow), Angioedema (broad), Asthma/bronchospasm (broad), Haematopoietic erythropenia (narrow), Haematopoietic leukopenia (narrow), Haematopoietic thrombocytopenia (narrow), Haemorrhage laboratory terms (broad), Interstitial lung disease (narrow), Neuroleptic malignant syndrome (broad), Systemic lupus erythematosus (broad), Anticholinergic syndrome (broad), Arrhythmia related

investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Retroperitoneal fibrosis (broad), Dementia (broad), Pseudomembranous colitis (broad), Embolic and thrombotic events, venous (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Accidents and injuries (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Other ischaemic heart disease (narrow), Noninfectious diarrhoea (narrow), Respiratory failure (narrow), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (narrow), Hypoglycaemia (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (narrow), Sepsis (narrow), Opportunistic infections (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 8 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zyprexa 15 mg tablet once daily Levothyroxine 100 mcg tablet once daily Metformin ER 500 mg tablet 2 tablets by mouth twice daily Divalproex DR 240 mg Give 3 tablets by mouth once daily Metoprolol tartrate 50 mg tablet by mouth twice daily
Current Illness: COVID 19 (tested positive on 11/27/2020). Appeared to have recovered, but had some weight loss as well as persistent weakness, activity intolerance.
Preexisting Conditions: Schizophrenia Dementia without behavioral disturbance Personal history of covid-19 (Tested positive on 11/27/2020) Type 2 diabetes mellitus Morbid Obesity Essential Hypertension BPH with LUTS Allergic rhinitis Hypothyroidism Seborrheic dermatitis Fatty liver disease Pancytopenia Glaucoma History of nicotine dependence, cigarettes
Allergies: Ciclopirox- erythema and pruritus Naltrexone- reaction dizziness Topiramate- mood changes
Diagnostic Lab Data: 12/31/2020: WBC 4.8 K/uL ; RBC 4.23 M/Ul; Hgb 12.9 g/dL; Hct 38.6%; Platelets 63 k/uL; Blood culture: No growth; Urine culture: Moderate mixed flora; Lipase 55 U/L; Procalcitonin 2.38 ng/mL; BNP 90; Lactic acid 7.2 mmo/L; CRP 110.5 mg/L; Potassium 3.4 mmol/L; INR 2.4 ; CT Angio Chest: Pulmonary embolism left and right; Moderate pachy peripheral ground glass right infiltrates Heavy triple vessel coronary calcification with heavy left main coronary calcification. Mild inflammatory stranding around the normal appearing pancreatic head suggesting pancreatitis. Moderate thickening of the wall of distal esophagus associate with small hiatal hernia: EKG on 12/31/2020: Sinus tachycardia; EKG on 1/2/2021: Atrial fibrillation with rapid RVR; On 1/3/2021: CT abdomen with contrast liver cirrhosis with portal venous hypertension, multiple splenic granulomata, recanalization of the umbilical vein. Third spacing with body wall edema and mild pericholecystic ascites. The mild stranding in the upper abdominal fat could represent mild pericholecystic ascites. Mostly liquid stool throughout colon without findings of intestinal obstruction.
CDC Split Type:

Write-up: On 12/31/2020, at approximately 00:15, pt developed a fever of 102.9 F and tachycardia with rate of 120. He was treated with acetaminophen. Later in the morning, he

complained of nausea, generalized muscle aches, intermittent increase in confusion. At approximately 14:00, he had a fall out of bed and at that time noted to be short of breath, tachypneic. He was taken via ambulance to Emergency Department. From there he was transferred to Hospital for admission with acute respiratory distress, suspected sepsis with lactic acid 7.4 and Bilateral Pulmonary Emboli. He was started on heparin and broad spectrum antibiotics and transitioned to ELIQUIS on 1/3/2021. Infectious etiology of sepsis was unclear. He continued broad spectrum antibiotics with clinical improvement. Abdominal CT scan was obtained due to intermittent nausea, vomiting, abdominal pain, loose stools. His heart rhythm flipped to Atrial Fibrillation with RVR on 1/2 and his rate improved with titration of metoprolol. He was also treated with prednisone for suspected underlying undiagnosed COPD. It is noted in his hospital summary that PEs presumed provoked in the setting of his recent COVID 19 infection. He was discharged from the hospital on 1/8/2021 and readmitted to the Veterans Home. He has been stable.

VAERS ID: 937582 (history)
Form: Version 2.0
Age: 38.0
Sex: Female
Location: Virginia
Vaccinated: 2021-01-05
Onset: 2021-01-08
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Dyspnoea, Influenza virus test negative, Laboratory test normal, SARS-CoV-2 test negative, Vomiting, White blood cell count increased
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Neuroleptic malignant syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: none
Preexisting Conditions: Asthma
Allergies: coconut

Diagnostic Lab Data: Evaluation performed by hospital-unsure of all testing completed but stated that she was negativeX3 for COVID, neg flu, and neg pneumonia.
CDC Split Type:

Write-up: Received Moderna vaccine on 1/5. Had a single episode of vomiting approximately 1 hour after vaccine. Hx of Asthma, started to develop SOB approximately 2 days later. She reached out to PCP and used inhaler with little relief. SOB worsened and she was admitted to the hospital on 1/8/21. Eval was negative for COVID (3 tests completed), flu and pneumonia. She has elevated WBCs and was given steroids and supplemental oxygen. She is improving but remains inpatient.

VAERS ID: 937774 (history)
Form: Version 2.0
Age: 30.0
Sex: Male
Location: Unknown
Vaccinated: 2021-01-05
Onset: 2021-01-10
Days after vaccination: 5
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / UNK	LA / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Cough, Pain, Pneumonia, Pyrexia, SARS-CoV-2 test positive, Sepsis

SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Sepsis (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Had Covid + PCR test on 10/06/2020.
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Fever/Cough/Body Aches. Covid +, Pneumonia and Sepsis

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VAERS ID: 937932 (history)
Form: Version 2.0
Age: 28.0
Sex: Male
Location: Utah
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Chest pain, Electrocardiogram ST segment elevation, Myalgia, Myocarditis, Pyrexia, Troponin increased
SMQs:, Rhabdomyolysis/myopathy (broad), Neuroleptic malignant syndrome (broad), Myocardial infarction (narrow), Anticholinergic syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: No known chronic conditions
Allergies: Spring seasonal allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient presented with myalgias, fevers, and chest pain on 1/10/21 and was found to have diffuse ST elevation and elevation troponin. He was evaluated by cardiology and diagnosed with acute myopericarditis. He was treated with NSAIDs and colchicine. He improved with this treatment and was discharged on 1/12/21 with ibuprofen and colchicine and outpatient cardiology follow up.

VAERS ID: 938067 (history)
Form: Version 2.0
Age: 95.0
Sex: Female
Location: Florida
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EH9899 / 2	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Cardio-respiratory arrest, Dyspnoea, Respiratory tract congestion, Tachycardia, Unresponsive to stimuli
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (narrow), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Respiratory failure (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cardiac murmur (Verbatim: Cardiac murmur); Congestive heart failure (Verbatim: Congestive heart failure); Hypertension (Verbatim: Hypertension)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021010600

Write-up: she was transferred to the hospital with tachycardia, congestion, shortness of breath, and is now unresponsive; she was transferred to the hospital with tachycardia, congestion, shortness of breath, and is now unresponsive; she was transferred to the hospital

with tachycardia, congestion, shortness of breath, and is now unresponsive; she was transferred to the hospital with tachycardia, congestion, shortness of breath, and is now unresponsive; This is a spontaneous report from a contactable nurse. A 95-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), Lot# EH9899 Exipration on 06Jan2021 at 15:00 at SINGLE DOSE at deltoid for COVID-19 immunization. The patient received first dose of the same vaccine BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), Lot# EH9899 Exipration date : 31Mar20221, on 16Dec2020. Medical history included : cardiac failure congestive, hypertension, cardiac murmur .There were no concomitant medications. The patient previously took cymbalta , vasotec and zocor and experienced drug hypersensitivity. The patient didn''t receive any other vaccines within 4 weeks prior to the COVID vaccine.This nurse,worker at a skilled nursing facility, reported that this patient with a history of heart failure, received her second dose of the Pfizer-BioNTech Covid-19 vaccine yesterday, 06Jan2021, at 3pm. At 7am on 07Jan2021 she was transferred to the hospital with tachycardia, congestion, shortness of breath, and is now unresponsive. Patient was stable prior to vaccination, but will now be transferred to hospice care. The nurse added the patient had the second COVID vaccine on 06Jan2021 yesterday and has now been transport to hospital due to a drastic decline after the shot. It was explained that this morning around 7 am she was transferred to the hospital. She was experiencing tachycardia, shortness of breath, and congestion. The events started this morning around 6-6:30am. The patient was admitted to the hospital. The shot was given at the facility. She received it at 3pm on 06Jan2021, First dose was on 16Dec2020.The caller relays she didn''t know how aggressive the hospital will be for the patient. She was a full code when left and now a DNR and is unresponsive. The patient will be going on hospice care. The causality was reported as related. The outcome of the events was not recovered.; Sender''s Comments: The Company cannot completely exclude the possible causality between the reported tachycardia, shortness of breath, congestion, unresponsive, and the administration of the COVID 19 vaccine, BNT162B2, based on the reasonable temporal association. The patient''s pre-existing medical condition of cardiac failure congestive, hypertension, cardiac murmur are confounding factors. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RA, IEC, as appropriate.

VAERS ID: 938091 (history)
Form: Version 2.0
Age: 50.0
Sex: Female
Location: Illinois
Vaccinated: 2020-12-18
Onset: 2020-12-18
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Blood calcium decreased, Blood magnesium decreased, Blood potassium

decreased, Feeling abnormal, Full blood count, Haemoglobin decreased, Hypocalcaemia, Hypokalaemia, Hypomagnesaemia, Metabolic function test, Muscle spasms, Tetany
SMQs:, Rhabdomyolysis/myopathy (broad), Haematopoietic erythropenia (broad), Haemorrhage laboratory terms (broad), Dementia (broad), Dystonia (broad), Chronic kidney disease (broad), Tumour lysis syndrome (broad), Hypokalaemia (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: CRESTOR; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Glucose tolerance impaired; Tachycardia; Tension headaches
Allergies:
Diagnostic Lab Data: Test Date: 20201218; Test Name: Calcium; Result Unstructured Data: Test Result:5.7; Test Date: 20210104; Test Name: Calcium; Result Unstructured Data: Test Result:9.5 (normal); Test Date: 20201218; Test Name: Magnesium; Result Unstructured Data: Test Result:1.2; Test Date: 20201221; Test Name: Magnesium; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201223; Test Name: Magnesium; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201226; Test Name: Magnesium; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201228; Test Name: Magnesium; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201230; Test Name: Magnesium; Result Unstructured Data: Test Result:Unknown results; Test Date: 20210104; Test Name: Magnesium; Result Unstructured Data: Test Result:2.1 (normal); Test Date: 20201218; Test Name: Potassium; Result Unstructured Data: Test Result:2.6; Test Date: 20210104; Test Name: Potassium; Result Unstructured Data: Test Result:4.2 (normal); Test Date: 20201218; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201221; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201223; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201226; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201228; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201230; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20210104; Test Name: CBC; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201218; Test Name: hemoglobin; Result Unstructured Data: Test Result:10.2; Comments: (at normal for her); Test Date: 20210104; Test Name: hemoglobin; Result Unstructured Data: Test Result:14.5 (normal); Test Date: 20201218; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201221; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201223; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201226; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201228; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results; Test Date: 20201230; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results; Test Date: 20210104; Test Name: CMP; Result Unstructured Data: Test Result:Unknown results
CDC Split Type: USPFIZER INC2021015699

Write-up: Hypokalemia; Hypomagnesemia; Hypocalcemia; Hemoglobin dropped little bit; Tetany; Muscle cramps; This is a spontaneous report from a contactable healthcare professional, a physician assistant. A 50-year-old female patient received the first dose of the BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Lot Number: UNKNOWN), via an unspecified route of administration on 18Dec2020 (at the age of 50-years-old) as a single dose for COVID-19 immunization. Medical history included tachycardia, tension headache, and glucose tolerance impaired. Concomitant medications included rosuvastatin calcium (CRESTOR), gabapentin (MANUFACTURER UNKNOWN), and metoprolol (MANUFACTURER UNKNOWN). On 18Dec2020, the patient experienced hypokalemia, hypomagnesemia, hypocalcemia, tetany, muscle cramps, and hemoglobin dropped little bit. The clinical course was as follows: The patient received the vaccine and about 30 minutes later she started to "feel bad". She went to the urgent care and about an hour after they took her to the emergency room (also reported as hospital). She had a complete blood count, complete metabolic panel, and magnesium level done on the 18th, 21th, 23rd, 26th, 28th, 30th of Dec2020 and then again on 04Jan2021. On 18Dec2020, her initial lab test showed potassium: 2.6, hemoglobin: 10.2, magnesium: 1.2, and calcium: 5.7. The physician assistant reported that a potassium of 2.6 was critically low and a hemoglobin was 10.2 was about normal for the patient. She received oral medications of potassium 20 mEq twice a day and calcium and magnesium supplements once a day. On 04Jan2021, lab data showed: potassium: 4.2, magnesium: 2.1, calcium: 9.5 and hemoglobin: 14.5. Per the reporting physician assistant, the above levels went back to normal but stated she had to be supplemented to get back to that point. The clinical outcomes of the hypokalemia, hypomagnesemia, hypocalcemia, and hemoglobin dropped little bit were recovered on 04Jan2021; while that of the tetany and muscle cramps, were unknown. The physician assistant assessed the events as related to the suspect vaccine. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.; Sender"s Comments: A possible contributory effect of suspect BNT162B2 on reported events cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.

VAERS ID: 938104 (history)
Form: Version 2.0
Age: 70.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-22
Onset: 2020-12-28
Days after vaccination: 6
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: SARS-CoV-2 test positive

SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: COVID-19 positive and admitted to hospital 6 days post-vaccination

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VAERS ID: 938109 (history)
Form: Version 2.0
Age: 43.0
Sex: Male
Location: Iowa
Vaccinated: 2020-12-31
Onset: 2021-01-01
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	O11J20A / 1	AR / IM

Administered by: Work **Purchased by:** ?
Symptoms: COVID-19 pneumonia, Chills, Cough, Headache, Pain, Pyrexia, SARS-CoV-2 test positive, Sinus congestion
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: COVID swab positive 1/1/2021 Hospital admission 1/11/2021 not sure what tests are being performed
CDC Split Type:

Write-up: Received first dose Moderna COVID vaccine on 12/28/2020; on that same day 12/28 he noticed a dry cough; on 1/1/2021 he reported fever, chills, body aches, Headache, sinus congestion. On 1/1/2021 he tested positive for COVID-19, he reported being with a family member on Christmas who had COVID symptoms. On 1/11/2021 he required hospitalization for COVID-19 pneumonia

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VAERS ID: 938118 (history)
Form: Version 2.0
Age: 51.0
Sex: Female
Location: Michigan
Vaccinated: 2021-01-05
Onset: 2021-01-08
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 1	- / IM

Administered by: Public **Purchased by:** ?
Symptoms: Abdominal X-ray, Aneurysm, Angiogram cerebral, Arteriogram carotid, Cerebellar haemorrhage, Cerebrovascular accident, Chest X-ray, Computerised tomogram abdomen, Computerised tomogram head, Computerised tomogram pelvis, Computerised tomogram spine, Computerised tomogram thorax
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow)

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 2021-01-10
Days after onset: 2
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: multivitamin; vitamin D
Current Illness: none mentioned
Preexisting Conditions: no significant past medical history
Allergies: latex
Diagnostic Lab Data: CT brain head; CT cervical spine; chest xray; CT chest, abdomen, pelvis; abdominal xray; CT angiogram neck & brain
CDC Split Type:

Write-up: on 1/8/2021 17:30 patient taken to ER, cerebellar hemorrhage, stroke, aneurysm

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VAERS ID: 938126 (history)
Form: Version 2.0
Age: 32.0
Sex: Male
Location: New York
Vaccinated: 2021-01-07
Onset: 2021-01-10
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK5730 / 1	- / -
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3246 / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Facial asymmetry, Facial paralysis, Fatigue, Headache, Myalgia, Paraesthesia
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious meningitis (broad), Eosinophilic pneumonia (broad), Hearing impairment (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: none

Current Illness: none
Preexisting Conditions: none
Allergies: none
Diagnostic Lab Data:
CDC Split Type:

Write-up: Following the first COVID vaccine dose on Dec/18/2020, I had headaches that started on the third day and ended on the tenth day. The headaches were usually light, unilateral, and alternating from one side to the other. I was usually functional except on the fourth and seventh days where the headaches were moderate to severe, and I took naps to help with the headaches for those two days. I have never had an issue with headaches before, and these symptoms were a new experience for me. I did not take any medications as treatment for the headaches. Following the second COVID vaccine dose on January/7/2021, I felt fatigue and generalized muscle aches within six to twelve hours, and these symptoms lasted for two days. On January/10/2021, when I woke up that morning I again felt light, unilateral, and alternating headaches. In addition, I noticed that I was unable to move the left side of my face. I felt moderate tingling sensations associated with the distribution of the paralysis. When I looked in the mirror, I could quite noticeably see asymmetry in my face. I immediately went to the emergency department at the hospital where my primary care doctor is located. I was kept in the hospital into the next day for observation. After evaluation by a neurology team and an MRI, I was provided with the diagnosis of Bells Palsy. I have never previously been diagnosed with Bells Palsy, and I have never previously had a hospital stay before. The doctors prescribed medications which I am currently taking. As of today January/12/2021, the symptoms have had some improvement, but the symptoms still continue.

VAERS ID: 938235 (history)
Form: Version 2.0
Age: 40.0
Sex: Male
Location: New Jersey
Vaccinated: 2021-01-06
Onset: 2021-01-08
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Blood test normal, Diarrhoea, Electrocardiogram normal, Hot flush, Nausea, Palpitations, Vomiting
SMQs:, Acute pancreatitis (broad), Arrhythmia related investigations, signs and symptoms (broad), Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Noninfectious diarrhoea (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes

Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None at the time of vaccination. Up to one month prior the individual had a cold described as coughing and sneezing
Preexisting Conditions: None
Allergies: None, possibly penicillin
Diagnostic Lab Data: Bloodwork and EKG at the hospital on 1/8 both of which were normal.
CDC Split Type:

Write-up: Beginning at 6:30 am on 1/8, two days following vaccination, the individual was driving to work and began to experience nausea while at a traffic light. He pulled over and threw up. Once he got to work he vomited again. He drove to another facility for work and then threw up 3 or 4 more times. He began experiencing hot flashes at this time. He went to the hospital and was seen in the emergency room. He threw up several more times in the ER and stated that his heart was racing and that he had diarrhea. The emergency room gave him medications for the nausea and he stayed there for 2 or 3 hours. He began to feel better and was discharged to home.

VAERS ID: 938256 (history)
Form: Version 2.0
Age: 53.0
Sex: Female
Location: New York
Vaccinated: 2021-01-06
Onset: 2021-01-06
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011L20A / 1	LA / IM

Administered by: Work **Purchased by:** ?
Symptoms: Blood test normal, Burning sensation, Chest X-ray normal, Dizziness, Electrocardiogram normal, Fatigue, Gait disturbance, Headache, Immediate post-injection reaction, Pain, Syncope, Ultrasound scan normal
SMQs:, Torsade de pointes/QT prolongation (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vyvanse 40mg daily took that morning. mvi and calcium 600mg OTC. Generic Voltaran 1% gel apply once a day.
Current Illness: none
Preexisting Conditions: Takes medications for weight loss Chronic neck and right shoulder pain
Allergies: Penicillin, erythromycin.
Diagnostic Lab Data: When to ER 1/8/2021- treated and release that day. Dx Syncope. States felt worse after discharge. ultrasound over injections site, blood work, ECG and CXR- everything normal. took Tylenol and Motrin.
CDC Split Type:

Write-up: Immediately developed intense burning that progressed over the next 30 minutes and continued to burn for 3 days. The next day progressive extreme fatigue last 4 days. felt like it was going to pass out. full on body pain, dizzy and lightheaded. need assistance to get to the walk, intense headache.

VAERS ID: 938258 (history)
Form: Version 2.0
Age: 48.0
Sex: Female
Location: Pennsylvania
Vaccinated: 2020-12-26
Onset: 2020-12-27
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / -

Administered by: Private **Purchased by:** ?
Symptoms: Injection site erythema, Injection site pain, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: Aspirin, barbiturates, NSAIDs
Diagnostic Lab Data:
CDC Split Type:

Write-up: Very large, reddened, tender area at injection site (like the size of an angry red egg), and fever of 103.2.

VAERS ID: 938361 (history)
Form: Version 2.0
Age: 53.0
Sex: Male
Location: Illinois
Vaccinated: 2021-01-06
Onset: 2021-01-10
Days after vaccination: 4
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	RA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Lip swelling, Urticaria
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Cetirizine 10mg - 1 daily, Fluoxetine 40mg - 1 daily, Jardiance 10mg - 1 Daily, Pazeo 0.7% eye drop - 1 drop into both eyes once daily, Famotidine 20mg - 1 twice daily, Fluticasone 50mcg nasal spray - 1 spray in each nostril twice daily, GI
Current Illness: Positive COVID 19 on 11/30/2020
Preexisting Conditions: Moderate Intellectually disabled, Non-Insulin Dependent Diabetes Mellitus, HTN, high cholesterol, schizophrenia, tachycardia, acid reflux
Allergies: Bee venom
Diagnostic Lab Data: Admitted to Hospital for Observation on 1/10/2021
CDC Split Type:

Write-up: Patient presented on the morning of 1/10/2021 with swollen lips and hives. Vaccination took place on Wednesday January 6th, 2021.

VAERS ID: 938397 (history)
Form: Version 2.0
Age: 57.0
Sex: Female
Location: Kansas
Vaccinated: 2021-01-04
Onset: 2021-01-11
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / IM

Administered by: Private **Purchased by:** ?
Symptoms: Arthralgia, C-reactive protein increased, Condition aggravated, Fibrin D dimer increased, Joint range of motion decreased, Neutrophil percentage increased, Pain, Platelet count increased, Red blood cell sedimentation rate increased
SMQs:, Haemorrhage laboratory terms (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: 1/11/21 CRP = 5.5. SED rate =73, D dimer= 1170, Platelet= 418, Neutrophils=78,
CDC Split Type:

Write-up: Patient had mild bilateral knee and hip pain 1 month ago, then she received her 1st dose of COVID vaccine on 12/14/20. Her joint pain worsened then on 1/4/21 she received her 2nd dose of COIVD vaccine on 1/4/21 and then her pain increased in spread up to her shoulders. She was seen at immediate care on 1/11/21 because her joint pain had worsened throughout and her shoulder pain was making it difficult to raise her arms above the level of

her shoulders. Her pain was made worse w/ movement.

VAERS ID: 938425 (history)
Form: Version 2.0
Age: 44.0
Sex: Female
Location: Texas
Vaccinated: 2020-12-30
Onset: 2021-01-06
Days after vaccination: 7
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Atrial fibrillation, Blood test, Dizziness, Echocardiogram, Electrocardiogram abnormal, Hot flush, Intensive care, Palpitations
SMQs:, Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Supraventricular tachyarrhythmias (narrow), Cardiomyopathy (broad), Vestibular disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Zyrtec
Current Illness: None
Preexisting Conditions: None
Allergies: Reglan, Bactrim, Augmentin
Diagnostic Lab Data: EKG, ECHO, blood work all done on 1/6/2021-1/8/2021
CDC Split Type:

Write-up: Woke up on 1/6/2021 with hot flashes, palpitations, dizziness and heart racing. Went to urgent care and they did an EKG which showed A-Fib, so I was sent to the ER and from there, I was transferred to an ICU at a different facility . I stayed until 1/8/2021. No cause was found and no history of A-Fib or family history.

VAERS ID: 938446 (history)
Form: Version 2.0
Age: 20.0

Sex: Female
Location: Iowa
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20-2A / 1	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Injection site erythema, Injection site swelling, Pyrexia
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Extravasation events (injections, infusions and implants) (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Fever (103-104 oF) and 4"x1" red, swelling area around site of injection. Pt states she received an IV.

VAERS ID: 938482 (history)
Form: Version 2.0
Age: 61.0
Sex: Unknown
Location: Unknown
Vaccinated: 2020-12-30
Onset: 2021-01-01
Days after vaccination: 2
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025J20A / 1	LA / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Acute respiratory failure, Chronic obstructive pulmonary disease
SMQs:, Anaphylactic reaction (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (broad), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (broad), Toxic-septic shock conditions (broad), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (narrow), Hypersensitivity (broad), Respiratory failure (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Acute approximate respiratory failure secondary to acute COPD

VAERS ID: 938524 (history)
Form: Version 2.0
Age: 43.0
Sex: Female
Location: New York
Vaccinated: 2021-01-08
Onset: 2021-01-11
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3246 / 2	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Anaphylactic reaction, Blood magnesium normal, Blood potassium decreased, Chest X-ray normal, Dyspnoea, Fibrin D dimer normal, Full blood count, Headache, Human chorionic gonadotropin negative, Intensive care, Metabolic function test abnormal, Nausea, Pain, Procalcitonin normal, Pruritus, Tachycardia, Urticaria, White blood cell count increased

SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Anaphylactic/anaphylactoid shock conditions (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), Hypokalaemia (narrow)

Life Threatening? Yes
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: mg, vit b complex, calicum with vit D, zyrtec, singulair, klonopin, omeprazole, prednisone
Current Illness: non systemic vasculitis bilat feet
Preexisting Conditions: asthma, seasonal allergies, feet non systemic vasculitis, colon polyps
Allergies: peanuts, tree nuts, compazine, ct dye
Diagnostic Lab Data: cbc, mild elevated wbc, cmp abn low K 2.8, d dimer neg, procalcitonin neg, hcg negative, chest xray negative, mg normal,
CDC Split Type:

Write-up: first day after shot, nausea, body aches, 2nd day Sunday headache, Monday 5 am woke up itching, then 9 am hives everywhere, trouble breathing, anaphylaxis, went to ER, got epi X 2, solumedrol, benadryl, pepcid, then still with hives, tachycardia, dyspnea, iv fluids were influsing and epi drip started, went to ICU

VAERS ID: 938576 (history)
Form: Version 2.0
Age: 22.0
Sex: Female
Location: New Jersey
Vaccinated: 2021-01-06
Onset: 2021-01-09
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 2	LA / SYR
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) /	EH9899 /	LA / SYR

PFIZER/BIONTECH	1	

Administered by: Private **Purchased by:** ?
Symptoms: Back pain, Blood test, Computerised tomogram, Deep vein thrombosis, Magnetic resonance imaging, Pulmonary embolism
SMQs:, Retroperitoneal fibrosis (broad), Embolic and thrombotic events, venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 2 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: None
Allergies: All seafood
Diagnostic Lab Data: CT, MRI, blood teatst
CDC Split Type:

Write-up: Back pain, bilateral PE and DVT

VAERS ID: 938712 (history)
Form: Version 2.0
Age: 36.0
Sex: Female
Location: Kansas
Vaccinated: 2021-01-06
Onset: 2021-01-07
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EK9231 / 2	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Blood glucose increased, Gait disturbance, Influenza like illness, Intervertebral disc protrusion, Magnetic resonance imaging spinal abnormal, Muscle fatigue, Muscular weakness, Paraesthesia, Red blood cell sedimentation rate increased, Respiratory rate
SMQs:, Rhabdomyolysis/myopathy (broad), Peripheral neuropathy (broad), Hyperglycaemia/new onset diabetes mellitus (narrow), Anticholinergic syndrome (broad), Parkinson-like events (broad), Guillain-Barre syndrome (broad), Noninfectious

encephalopathy/delirium (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Metformin, insulin, lisinopril, omeprazole, atorvastatin, propranolol
Current Illness: DM type II insulin dependent, hypertension, hyperlipidemia
Preexisting Conditions: Reactive airway disease, diabetes mellitus type II, PCOS (polycystic ovarian syndrome), muscle weakness postpartum course of weeks and months. Mother has an autoimmune disease.
Allergies: codeine - hypersensitivity, diphenhydramine - hypersensitivity, iodinated contrast - nausea, trouble breathing, clindamycin - hives
Diagnostic Lab Data: Abnormal labs as follows; blood glucose 206, ESR Westergren 26, MRI of lumbar spine - only abnormal finding was bulging L5S1. Vitals normal except for respiratory rate
CDC Split Type:

Write-up: Patient states that she received her second vaccination and in the hours after she had flu-like symptoms. Then over the next few days, she started to notice tingling and a "prickly" sensation in various areas. This progressed to symmetric BLE weakness which started in her feet and had reached to just above her knees bilaterally time and she arrived. The weakness had progressed to her hips. She also noticed weakness in her arms and they are easily fatigued. She is able to walk but it takes much effort.

VAERS ID: 938808 (history)
Form: Version 2.0
Age: 52.0
Sex: Female
Location: Unknown
Vaccinated: 2021-01-09
Onset: 2021-01-09
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037K20A / 1	- / -

Administered by: Private **Purchased by:** ?
Symptoms: Cough, Cyanosis, Flushing, Hyperhidrosis
SMQs:, Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Acute central respiratory depression (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: codeine, Combivent
Current Illness:
Preexisting Conditions: Behcet''s syndrome, asthma, laryngeal spasms, Guillain-Barre, Raynaud''s, GERD, anxiety/depression
Allergies: quinolones, contrast media, linezolid, vancomycin, sulfa, shellfish, daptomycin, tocilizumab
Diagnostic Lab Data:
CDC Split Type:

Write-up: coughing, flushing, cyanosis, diaphoresis

VAERS ID: 938880 (history)
Form: Version 2.0
Age: 57.0
Sex: Male
Location: Virginia
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Brain natriuretic peptide increased, C-reactive protein increased, COVID-19 pneumonia, Cardiomegaly, Chest X-ray abnormal, Chest discomfort, Computerised tomogram thorax, Dyspnoea, Fatigue, Fibrin D dimer increased, Haematocrit decreased, Haemoglobin decreased, Impaired work ability, Liver function test, Lung opacity, Metabolic function test, Neutrophil count increased, Pain, Pericardial effusion, Renal function test, Respiratory viral panel, SARS-CoV-2 test positive, Scan with contrast abnormal, Troponin I normal, Urine analysis, White blood cell count normal
SMQs:, Cardiac failure (broad), Anaphylactic reaction (broad), Haematopoietic erythropenia (broad), Haemorrhage laboratory terms (broad), Interstitial lung disease (narrow), Systemic lupus erythematosus (broad), Malignancy related therapeutic and diagnostic procedures (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Drug reaction with eosinophilia and systemic symptoms

syndrome (broad), Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Prednisone 2mg every other day (Cut from 3mg on 12/16/20) Te Prednisone 2mg every other day (Cut from 3mg on 12/16/20) Telmisartan 80mg daily Norvasc 10mg daily Coreg 25mg BID Bumex 1mg po daily PRN ? not needed recently Aspar
Current Illness: denies
Preexisting Conditions: G6PD positive, HTN, DM, prostate CA, MCD ? steroid dependent, chronic renal disease
Allergies: Meloxicam
Diagnostic Lab Data: 1/11/2021: CXR = There is scattered nodular ground glass and confluent opacification of the lungs bilaterally, primarily within the posterior regions and lower lobes. No evidence of 1/11/2021: CXR = There is scattered nodular ground glass and confluent opacification of the lungs bilaterally, primarily within the posterior regions and lower lobes. No evidence of pleural effusion or pneumothorax. Findings: no PE, trace pericardial effusion with mild generalized cardiomegaly; atypical infectious vs inflammatory etiology, such as covid 19 pneumonia. Cta pulmonary arteries w/wo contrast: no PE, same as cxr for effusion, pneumonia suggestion. 1/11/2021 Labs: COVID 19 positive, D-Dimer 0.77 (0.27-0.50), CRP 156.2 (<9.9), RVP panel was all negative; BNP 47, WBC 7.5 with 90 neutrophils, h/h 11.1/34.6, troponin I <0.012, and has lots of other labs ? CMP, HFT, renal panels, UA, etc
CDC Split Type:

Write-up: this 57yo Male ret. He received his first Pfizer covid vaccine on Jan 7th to left arm at hospital (not where he works). He went home and reported was fine, but tired. Stated his fatigue and onset of shortness of breath began day 1 post vaccine kept him from working. As his symptoms worsened, on day 4 (1/11/2021) he obtained a pulse ox and O2 sats were in the "80"s" with normal heart rate. He went to ER on 1/11 1507 with body aches, fatigue, chest discomfort and SOB where he was ultimately admitted with diagnosis of COVID-19 pneumonia. Is on O2 support via NC.

VAERS ID:	938958 (history)
Form:	Version 2.0
Age:	30.0
Sex:	Female
Location:	Illinois
Vaccinated:	2021-01-12
Onset:	2021-01-12
Days after vaccination:	0
Submitted:	0000-00-00

Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Immediate post-injection reaction
SMQs:, Anticholinergic syndrome (broad), Vestibular disorders (broad), Hypersensitivity (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Vitamins
Current Illness: no
Preexisting Conditions: no
Allergies: no
Diagnostic Lab Data: no
CDC Split Type:

Write-up: Immediately after she felt faint, heart rate 121, felt faint again bp 62/33. Was taken to the ER, within a half hour, she fully recovered. Vitals went back to normal.

VAERS ID: 939006 (history)
Form: Version 2.0
Age: 45.0
Sex: Female
Location: Illinois
Vaccinated: 2021-01-12
Onset: 2021-01-12
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Dizziness, Hypoaesthesia, Pain in extremity, Peripheral coldness
SMQs:, Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Vestibular disorders (broad), Tendinopathies and ligament disorders (broad), Sexual dysfunction (broad)

Life Threatening? No
Birth Defect? No

Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations: reaction to HEP B
Other Medications: Vitamins
Current Illness: no
Preexisting Conditions: no
Allergies: Hep B
Diagnostic Lab Data: no
CDC Split Type:

Write-up: Severely dizzy, left hand totally numb but painful, cold to touch. Felt better before she got to ER.

VAERS ID: 939087 (history)
Form: Version 2.0
Age: 67.0
Sex: Female
Location: Indiana
Vaccinated: 2021-01-07
Onset: 2021-01-08
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / UNK	LA / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Blood culture negative, Body temperature increased, CSF test normal, Mental status changes, Urine analysis normal, White blood cell count normal
SMQs:, Neuroleptic malignant syndrome (broad), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 5 days
Extended hospital stay? No
Previous Vaccinations:

Other Medications: Amlodipine, Cholecalciferol, Coumadin, Pantoprazole
Current Illness: Was staying at Rehab Center after having hip surgery after 12/20/2020 but unsure of exact date of surgery.
Preexisting Conditions: Mass on brain, TIAs, Craniotomy 8/2018, HTN, cholecystectomy, appendectomy, tubal ligation, R total hip replacement, Hepatitis years ago, cancer of brain and lung, IVC filter, tonsillectomy
Allergies: NKA
Diagnostic Lab Data: She had a temp of 102 and altered mental status. She had a full septic workup including CSF. It was all neg. Normal WBC, urine, blood cultures and CSF. Normalized and sent back to Rehab.
CDC Split Type:

Write-up: Attempting to confirm which COVID 19 vaccine was given (Moderna or Pfizer). They did not send the record when they sent the patient to General ER the next am. Did not answer the phone.

VAERS ID: 939564 (history)
Form: Version 2.0
Age: 29.0
Sex: Female
Location: Ohio
Vaccinated: 2020-12-22
Onset: 2021-01-05
Days after vaccination: 14
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain upper, Computerised tomogram abdomen abnormal, Computerised tomogram pelvis abnormal, Lipase increased, Magnetic resonance cholangiopancreatography, Nausea, Pancreatitis acute
SMQs:, Acute pancreatitis (narrow), Retroperitoneal fibrosis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 3 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Lexapro 20mg daily, Wellbutrin 300mg daily, Zyrtec OTC
Current Illness: None
Preexisting Conditions:

Allergies: Peanuts
Diagnostic Lab Data: Lipase on 1/5/2020 was $g1800 CT on 1/5/2020 of the abdomen/pelvis showed pancreatitis MRCP on 1/5/2020 was negative
CDC Split Type:

Write-up: On 1/5/2020, I, the patient woke up at 3:30 with sharp boring epigastric pain. Progressed to include nausea. On evaluation in the ED was diagnosed with acute pancreatitis. Prior hx includes 2 episodes with hospitalization for gallstone pancreatitis in 2016. Subsequently had laparoscopic cholecystectomy in 2016. Last alcoholic beverage prior to 2020 ED presentation was 3 weeks prior. No tobacco or drug use.

VAERS ID: 963164 (history)
Form: Version 2.0
Age: 49.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Cardiac failure congestive, Diabetes mellitus, Hyperglycaemia, Hypertension, Pyrexia, Tachycardia, Tachypnoea
SMQs:, Cardiac failure (narrow), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Hypertension (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type:

Write-up: Fever, Tachypnea, HYPERtension, Tachycardia & HYPERglycemia Narrative: Was inpatient overnight in a telemetry until. DC with diagnosis of DM and CHF

VAERS ID: 1402179 (history)
Form: Version 2.0
Age: 49.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-28
Onset: 2020-12-29
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Cardiac failure chronic, Cardiac telemetry, Diabetes mellitus, Hyperglycaemia, Hypertension, Pyrexia, Tachycardia, Tachypnoea
SMQs:, Cardiac failure (narrow), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Hypertension (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Dehydration (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Fever, Tachypnea, HYPERtension, Tachycardia & HYPERglycemia Narrative: Was inpatient overnight in a telemetry until. DC with diagnosis of DM and CHF

VAERS ID: 936161 (history)
Form: Version 2.0
Age: 28.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-27
Onset: 2020-12-27
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6796 / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Blood test, Hypersensitivity, Malaise, Rash pruritic, Tryptase, Vision blurred
SMQs:, Anaphylactic reaction (broad), Angioedema (broad), Anticholinergic syndrome (broad), Glaucoma (broad), Lens disorders (broad), Retinal disorders (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201227; Test Name: Blood test; Result Unstructured Data: Test Result:Nothing abnormal; Test Date: 20201227; Test Name: Blood tryptase; Result Unstructured Data: Test Result:Unknown results
CDC Split Type: DKPFIZER INC2021013306

Write-up: Assessed as an allergic reaction but not as regular anaphylaxis by the attending physician.; Itchy rash on thorax, neck and upper arms; Accompanied by slight general malaise; Accompanied by slightly blurred vision; This is a spontaneous report from a contactable physician via The Medicines Agency (MA) downloaded from the Medicines Agency (MA) Regulatory Authority-WEB DK-DKMA-WBS-0028060. A 28-years-old female patient received the first dose of bnt162b2 (COMIRNATY, lot: EJ6796, exp date 30Apr2021), intramuscular on 27Dec2020 at single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. On 27Dec2020, 13-15 minutes after the vaccination the patient developed an allergic reaction (Assessed as an allergic reaction

but not as regular anaphylaxis by the attending physician) with itchy rash, blurred vision and malaise. The outcome of the events was recovered on the same day, after few hours. The ADRs were by the physician reported as resulting in hospitalisation (27Dec2020, observed a few hours afterwards). The patient was treated with i.v methylprednisolone (SOLU-MEDROL) and clemastine (TAVEGYL). The patient underwent lab tests and procedures on 27Dec2020 which included blood test: nothing abnormal and tryptase: unknown results. The patient did not experience: No regular respiratory or circulatory reaction. The patient was ABC stable. The 2nd dose of the vaccine is planned given during antihistamine aler). If the Medicines Agency receives supplemental significant information regarding this case the case will be re-submitted.

VAERS ID: 936162 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-29
Onset: 2020-12-29
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6796 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Blood pressure measurement, Chills, Nausea, Paraesthesia, Presyncope
SMQs:, Acute pancreatitis (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Guillain-Barre syndrome (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201229; Test Name: Blood pressure; Result Unstructured Data: Test Result:stable; Comments: stable

CDC Split Type: DKPFIZER INC2021013308

Write-up: Interpreted as a vasovagal reaction; paraesthesia in the fingers; Chills; Nausea; This is a spontaneous report from a contactable physician downloaded from the Regulatory Authority-WEB DK-DKMA-WBS-0028075, Safety Report Unique Identifier DK-DKMA-ADR 24548356. A 37-year-old female patient received bnt162b2 (COMIRNATY, lot: EJ6796, exp.30Apr2021), intramuscular on 29Dec2020 at single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. On 29Dec2020 five min after the vaccination the patient experienced interpreted as a vasovagal reaction, paraesthesia in the fingers, chills, nausea. The events were serious for hospitalization at 29Dec2020. The patient underwent lab tests and procedures which included blood pressure measurement: stable on 29Dec2020. The outcome of the events was not resolved. No follow up attempts are possible. No further information is expected.

VAERS ID: 936164 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-16
Onset: 2020-12-16
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Angiogram, Chest pain, Electrocardiogram, Myocardial infarction, SARS-CoV-2 test, Troponin
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; ; ; SYMBICORT; ;
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: DVT; Pulmonary embolism
Allergies:

Diagnostic Lab Data: Test Name: Angiogram; Result Unstructured Data: Test Result:unknown; Test Name: Electrocardiogram; Result Unstructured Data: Test Result:unknown; Test Date: 20201216; Test Name: COVID-19 virus test; Result Unstructured Data: Test Result:Negative; Comments: No - Negative COVID-19 test; Test Name: Troponin; Result Unstructured Data: Test Result:unknown
CDC Split Type: GBPFIZER INC2021003917

Write-up: Chest pain; Myocardial infarct; This is a spontaneous report from a contactable pharmacist downloaded from the Agency Regulatory Authority-WEB. The regulatory authority report number is GB-MHRA-ADR 24545531. An 82-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 16Dec2020, at 17:45 at single dose for immunization. Medical history included pulmonary embolism and deep vein thrombosis, both from an unknown date. Concomitant medication included warfarin (MANUFACTURER UNKNOWN), paracetamol (MANUFACTURER UNKNOWN), salbutamol (MANUFACTURER UNKNOWN), codeine (MANUFACTURER UNKNOWN), budesonide, formoterol fumarate (SYMBICORT), aclidinium (MANUFACTURER UNKNOWN), amitriptyline (MANUFACTURER UNKNOWN). The patient experienced myocardial infarct on 16Dec2020 and chest pain on 16Dec2020, at 19:45. The events were serious as they were medically significant and involved hospitalization. The patient underwent lab tests and procedures which included angiogram and electrocardiogram on an unknown date with unknown outcome, COVID-19 virus test was negative on 16Dec2020. Details were as follows: patient had the vaccine at 17:45, then at 19:45, she had sudden chest pain; there was a call to emergency and was she was taken to accident and emergency at 22:04. No previous history of heart conditions was noted. Patient has not tested positive for COVID-19 since having the vaccine. The outcome of myocardial infarct was recovering; the outcome of chest pain was unknown. No follow-up attempts are possible, information on batch number cannot be obtained.

VAERS ID: 937983 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Foreign
Vaccinated: 2020-12-23
Onset: 2020-12-26
Days after vaccination: 3
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Body temperature, Hepatic enzyme, Hepatic enzyme increased, Joint range of motion decreased, Malaise, Mobility decreased, Myalgia, Pyrexia, Splenomegaly, Ultrasound abdomen
SMQs:, Rhabdomyolysis/myopathy (broad), Liver related investigations, signs and symptoms (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Parkinson-like events (broad), Eosinophilic pneumonia (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic

symptoms syndrome (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20201226; Test Name: temperature; Result Unstructured Data: Test Result:39 to 40 degrees; Test Name: hepatic enzymes; Result Unstructured Data: Test Result:moderate elevation; Test Name: abdominal ultrasound; Result Unstructured Data: Test Result:splenomegaly
CDC Split Type: CRPFIZER INC2021010005

Write-up: moderate elevation of hepatic enzymes; splenomegaly; limitation for mobilization of both shoulders more on the left side; severe generalized myalgia; limitation to move; fever of 39 to 40 degrees; general malaise; This is a spontaneous report from a contactable physician via a sales representative. A 69-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 23Dec2020 at single dose for Covid 19 Immunization. The patient medical history and patient"s concomitant medications were not reported. Patient received COVID 19 PFIZER vaccination on 23Dec2020. The next day(24Dec2020) he flew to a specific location and 48 hours later(26Dec2020) he started with severe generalized myalgia, fever of 39 to 40 degrees and general malaise. Initially in the 4 extremities and later more in the upper limbs, more in upper left limb than the right, he has limitation for mobilization of both shoulders more on the left side. He consulted on three occasions in consecutive days to emergencies of the Hospital due to limitation to move due to myalgia. The patient had moderate elevation of hepatic enzymes and splenomegaly was detected in the abdominal ultrasound. 24 hours ago(04Jan2021) he was hospitalized and started with Solumedrol at a dose of 40mg IV every 12 hours with significant improvement but he still had moderate pain and limited movement of the shoulders. Treatment was received for severe generalized myalgia and limitation to move, limitation for mobilization of both shoulders more on the left side. He was discharged with decreasing doses of Prednisone. The patient was hospitalized for moderate elevation of hepatic enzymes and splenomegaly, severe generalized myalgia, limitation to move, mobilization of both shoulders more on the left side, from 04Jan2021 to Jan2021. The outcome of events was unknown. In this patient, other possibilities of bacterial and viral infections were ruled out and we could not find anything to explain their manifestations. The report is made in order to report the manifestations of patient as possibly associated with recent vaccination. Information on the lot/batch number has been requested.; Sender"s Comments: Based on the compatible temporal association and in absence of strong confounding factors, the vaccination with BNT162B2 might play a contributory role in triggering the onset of the reported events. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics

Committees and Investigators, as appropriate.

VAERS ID: 937987 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-28
Onset: 2020-12-28
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6796 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Chest discomfort, Dysphagia, Nausea, Paraesthesia
SMQs:, Anaphylactic reaction (broad), Acute pancreatitis (broad), Peripheral neuropathy (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Depression
Allergies:
Diagnostic Lab Data:
CDC Split Type: DKPFIZER INC2021013452

Write-up: Paraesthesia in hands and feets; Nausea; Swallowing difficult; Chest pressure; This is a spontaneous report downloaded from the Agency Regulatory Authority-WEB DK-DKMA-WBS-0028054. The report was received from a contactable physician via Agency, regulatory authority number is DK-DKMA-ADR 24547917. A 55-year-old female patient received bnt162b2 (COMIRNATY) lot EJ6796, Exp date 30Apr2021, intramuscular on 28Dec2020 at single dose for covid-19 immunisation. Medical history included depression. Concomitant medication included sertraline (manufacturer unknown) for depression started in Oct2020. On 28Dec2020, the patient experienced paraesthesia in hands and feets, nausea,

swallowing difficult, chest pressure immediately after the injection. The events required hospitalization on 28Dec2020. The patient was treated with 0.3 mg adrenaline on the vaccination center. At the emergency department the patient received 1 mg clemastin and 50 mg prednisolone (per oral). The events recovered on 28Dec2020.

VAERS ID: 937988 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6797 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Allergy test, Blood test, Dizziness, Heart rate, Malaise, Palpitations, Presyncope, Tachycardia, Urticaria
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad), Vestibular disorders (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Food allergy ((almonds, green apples and carrots))
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Allergy test; Result Unstructured Data: Test Result:to be performed; Test Date: 20201230; Test Name: Blood test; Result Unstructured Data: Test Result:unspecified results; Test Date: 20201230; Test Name: Pulse rate; Result Unstructured Data: Test Result:115
CDC Split Type: DKPFIZER INC2021013364

Write-up: palpitation; dizzy; malaise; tachycardia; urticaria/rash; near fainting; This is a spontaneous report from a contactable physician downloaded from the Regulatory Authority-WEB DK-DKMA-WBS-0028102. The report was received via Agency, additional number DK-DKMA-ADR 24549048. A 29-year-old female patient on 30Dec2020 received bnt162b2 (COMIRNATY) (Batch/Lot No.: EJ6797, Exp date: 30Apr2021) intramuscular at single dose for covid-19 immunisation. Medical history included ongoing food allergy to almonds, green apples and carrots. There is no information regarding past medication. The patient''s concomitant medications were not reported. On 30Dec2020 after 20 minutes from vaccination the patient experienced palpitation, dizzy, malaise, tachycardia, urticaria/rash, near fainting. The patient did not experience swelling of the throat, dyspnea or stridor. The events resulted in hospitalisation on 30Dec2020. The patient underwent lab tests and procedures which included allergy test: to be performed on an unspecified date, blood test: unspecified results on 30Dec2020, Pulse rate: 115 on 30Dec2020. The outcome of the event urticaria/rash was resolving. The patient recovered from the events palpitation, dizzy, malaise, tachycardia, near fainting on 30Dec2020.

VAERS ID: 937989 (history)
Form: Version 2.0
Age: 32.0
Sex: Female
Location: Foreign
Vaccinated: 2020-12-30
Onset: 2020-12-30
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6797 / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Dysphonia, Headache, Pharyngeal swelling, Physical examination, Swollen tongue
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Parkinson-like events (broad), Oropharyngeal allergic conditions (narrow), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:

Current Illness: Allergy (Newly discovered allergy to rubber chemicals.); Anxiety (Several years)
Preexisting Conditions: Medical History/Concurrent Conditions: Rubber sensitivity; Comments: .
Allergies:
Diagnostic Lab Data: Test Date: 20201230; Test Name: Physical examination; Result Unstructured Data: Test Result:listened to the lungs and heart (no result describ; Comments: listened to the lungs and heart (no result described), looked in the mouth (nothing to see)
CDC Split Type: DKPFIZER INC2021013320

Write-up: Swelling in the throat /the throat began to narrow; the tongue swelled up; Headache; the voice was distorted; This is a spontaneous report from a contactable consumer (patient, Medical Laboratory Technologist) downloaded from the regulatory authority-WEB (Regulatory Authority number DK-DKMA-WBS-0028118, Safety Report Unique Identifier DK-DKMA-ADR 24549391). A 32-year-old female patient received the first single dose of BNT162B2 (COMIRNATY; lot/batch EJ6797, expiry date 30Apr2021) on 30Dec2020, for COVID-19 immunisation. Medical history included ongoing anxiety from several years and ongoing allergy from 2020 (newly discovered allergy to rubber chemicals). The patient''s concomitant medications were not reported. On 30Dec2020 the patient experienced swelling in the throat /the throat began to narrow, the tongue swelled up, headache, the voice was distorted. The patient was treated with two different agents in drip (not specified). All these events required hospitalization and resolved on 30Dec2020. On 30Dec2020 physical examination showed: listened to the lungs and heart (no result described), looked in the mouth (nothing to see). Allergy test was planned in 2021. If the Agency will receive supplemental significant information regarding this case the case will be re-submitted.

VAERS ID: 937991 (history)
Form: Version 2.0
Age: 22.0
Sex: Female
Location: Foreign
Vaccinated: 2021-01-02
Onset: 2021-01-02
Days after vaccination: 0
Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ6797 / 1	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: C-reactive protein, Flushing, Haematology test, International normalised ratio, Liver function test, Renal function test, Swelling face, Tryptase, Vaccination site erythema, Vaccination site swelling, White blood cell count
SMQs:, Anaphylactic reaction (broad), Angioedema (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 20210103; Test Name: C-reactive protein; Result Unstructured Data: Test Result:9,5; Test Date: 20210103; Test Name: Haematology test; Result Unstructured Data: Test Result:Normal; Test Date: 20210103; Test Name: International normalised ratio; Result Unstructured Data: Test Result:Normal; Test Date: 20210103; Test Name: Liver function test; Result Unstructured Data: Test Result:Normal; Test Date: 20210103; Test Name: Renal function test; Result Unstructured Data: Test Result:Normal; Test Date: 20210103; Test Name: Blood tryptase; Result Unstructured Data: Test Result:Unknown results; Test Date: 20210103; Test Name: Leucocyte count; Result Unstructured Data: Test Result:9,11
CDC Split Type: DKPFIZER INC2021013385

Write-up: Face swelling; Face flushing; Redness at the vaccination site; Swelling at the vaccination site; This is a spontaneous report from a contactable physician via The Agency downloaded from the Agency Regulatory Authority-WEB DK-DKMA-WBS-0028154, DK-DKMA-ADR 24551562. A 22-years-old female patient received the first dose of bnt162b2 (COMIRNATY, lot # EJ6797, exp date 30Apr2021) intramuscular on 02Jan2021 at single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Immediately after the vaccination on 02Jan2021 the patient developed vaccination site erythema and vaccination site swelling. The next morning on 03Jan2021 the patient wakes up early after bad sleep with facial flushing and facial swelling. The patient did not experience affected airways, no other kind of rash. The ADRs were by the reporter reported as resulting in hospitalisation on 03Jan2021. No treatment for the events was performed. The facial flushing and facial swelling were recovered on 03Jan2021. The vaccination site erythema and vaccination site swelling were recovering. The patient underwent lab tests and procedures on 03Jan2021 which included c-reactive protein: 9,5 unit not specified, haematology test: normal, international normalised ratio: normal, liver function test: normal, renal function test: normal, tryptase: Unknown results, white blood cell count: 9,11 unit not specified. If the Agency receives supplemental significant information regarding this case the case will be re-submitted.

VAERS ID: 938002 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2020-12-20
Onset: 2020-12-21
Days after vaccination: 1

Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EJ1688 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Dyspnoea, Nausea, Rash pruritic
SMQs:, Anaphylactic reaction (narrow), Acute pancreatitis (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: ; ; TRULICITY; ; ; BRALTUS
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: GBPFIZER INC2021008978

Write-up: Itchy rash, norsea , breathless; Itchy rash, norsea , breathless; Itchy rash, norsea , breathless; This is a spontaneous report from a contactable other hcp downloaded from the database GB-MHRA-EYC 00236074 Safety Report Unique Identifier GB-MHRA-ADR 24546104. A 57-year-old female patient started to receive bnt162b2, lot: Ej1688 intramuscularly on 20Dec2020 at single dose for COVID-19 immunisation. The patient medical history was not reported. Concomitant medication included metformin, risperidone, dulaglutide (TRULICITY), simvastatin, dapagliflozin, tiotropium bromide (BRALTUS). All concomitants taken for an unspecified indication from an unspecified date to an unspecified date. The patient experienced itchy rash, norsea (nausea), breathless on 21Dec2020 with seriousness criteria of hospitalization. The outcome of events was not recovered. No follow-up attempts possible. No further information expected.

VAERS ID: 938044 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Foreign
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Epilepsy
SMQs:, Systemic lupus erythematosus (broad), Convulsions (narrow), Generalised convulsive seizures following immunisation (narrow)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Seizures
Allergies:
Diagnostic Lab Data:
CDC Split Type: MXPFIZER INC2021013162

Write-up: epileptic state; This is a spontaneous report from a non-contactable consumer received from a Pfizer colleague. The reporter informed similar adverse events for three different patients. This is the third of three reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. The patient had a background of treated seizures crisis. The patient"s concomitant medications were not reported. The patient experienced epileptic state started 10 minutes after the vaccination on an unspecified date with outcome of unknown. The patient was hospitalized in an epileptic state. No follow-up attempts are possible; information about batch/lot number cannot be obtained.; Sender"s Comments: Linked Report(s) : MX-PFIZER INC-2021013153 same reporter/drug, different patient, similar event;MX-PFIZER INC-2021013154 Same reporter/drug, different patient, similar event

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VAERS ID: 938045 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Foreign
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1

Submitted: 0000-00-00
Entered: 2021-01-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Arthralgia, Dysphagia, Headache, Myalgia
SMQs:, Rhabdomyolysis/myopathy (broad), Anticholinergic syndrome (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Eosinophilic pneumonia (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, ? days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: ROPFIZER INC2021010266

Write-up: headache; myalgias; arthralgias; dysphagia; This is a spontaneous report from a nurse via a sales representative. A 36-year-old female patient received bnt162b2 (COMIRNATY, lot number and expiration date not reported) at single dose on 05Jan2021 10:00 a.m. for covid-19 immunization, administered at the Vaccination Center local city. The patient medical history and concomitant medications were not reported. On 6.01.2021 at 15:00 she accused the occurrence of headache, myalgias, arthralgias and dysphagia. She was hospitalized for supervision at local Hospital, but after 5 hours from hospitalization she requested discharge on demand, her condition being improved. The outcome of events was unknown. The information about lot number and expiration date has been requested.; Sender''s Comments: The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Based on the information available, a possible contributory role of the suspect products cannot be excluded for the reported events due to temporal association. Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators as appropriate.

VAERS ID: 939629 (history)

Form: Version 2.0
Age: 79.0
Sex: Female
Location: Louisiana
Vaccinated: 2021-01-05
Onset: 2021-01-06
Days after vaccination: 1
Submitted: 0000-00-00
Entered: 2021-01-13

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	L27L20A / 1	RA / SYR

Administered by: Private **Purchased by:** ?
Symptoms: Asthenia, Blood creatinine increased, Blood glucose normal, Blood potassium increased, Blood urine present, Confusional state, Culture urine, Diarrhoea, Fatigue, Haematocrit decreased, Haemoglobin decreased, Nausea, Pollakiuria, Pyrexia, Urinary incontinence, Vomiting
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (broad), Acute pancreatitis (broad), Haematopoietic erythropenia (broad), Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhage laboratory terms (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Dementia (broad), Pseudomembranous colitis (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Chronic kidney disease (broad), Noninfectious diarrhoea (narrow), Tumour lysis syndrome (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? No
Permanent Disability? No
Recovered? Yes
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? Yes, 1 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications: Amlodipine 10mg/day, Eliquis 2.5mg/day, Cyanocobalamine 1000mg Gabapentin 300/day, Lisinopril, 40mg/day, Metformin 500mg XR/day, Lopressor 25mg/day, Zofran 4mg/day, Protonix 40mg/day
Current Illness: Anticoagulant use, arthritis, asthma, AFib, bacteremia, CHG, lLumbar fractures, COPD, DM, DVT, Abcess, lung CA, Lupus, MRSA, Osteoporosis, PVD, Psoriasis, Sjorgrens, stroke, venous insufficiency of leg
Preexisting Conditions: See extensive list
Allergies: Bactrim, Codeine, Methotrexate, PCN, Sulfamethox
Diagnostic Lab Data: Hgb 9.1, HCT 29.8, Gluc 113, K+ 5.4, Cr. 2.0, urine culture taken
CDC Split Type:

Write-up: The patient presented to hospital on 1/6/2021 with a primary complaint of Fatigue (pt had covid vaccine yesterday. Now displaying increased weakness, blood in urine, increased confusion and urinated on herself today. Fever of 101.9) 79-year-old female

presents to the emergency room with fatigue. The patient states yesterday she had the Moderna COVID-19 The patient states today she had a temperature 101.8? prior to arrival. Her son noted that she was having episodes of urinary frequency and also that she was profoundly weak and fatigued. They denied falling or hitting her head. The patient also states she been having nausea vomiting with diarrhea.

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