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

Found 19,532 cases where Vaccine targets COVID-19 (COVID19) and Patient Died

Government Disclaimer on use of this data

Table

Age	Count	Percent
< 3 Years	4	0.02%
3-6 Years	1	0.01%
9-12 Years	1	0.01%
12-17 Years	34	0.17%
17-44 Years	737	3.77%
44-65 Years	2,322	11.89%
65-75 Years	2,752	14.09%
75+ Years	6,365	32.59%
Unknown	7,316	37.46%
TOTAL	19,532	100%

Case Details (Sorted by Onset Date)

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VAERS ID: 925264 (history)
Form: Version 2.0
Age: 77.0
Sex: Male
Location: Oklahoma
Vaccinated: 2020-12-31
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-07
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Norco 5/325 1 tablet every 6 hours as needed. Warfarin 4mg , 2 tablets orally Monday and Friday Torsemide 100mg, 1/2 tablet orally once a day Sabcubitril-Valsartan 24/26mg 1/4 tablet orally twice a day Gabapentin 300mg, tablet once daily Ca
Current Illness: CHF HTN DM 2 CKD Obesity DDD
Preexisting Conditions: CHF HTN DM 2 CKD Obesity DDD
Allergies: NKDA
Diagnostic Lab Data: None
CDC Split Type:

Write-up: PT was found deceased in his home on 1/5/2021

VAERS ID: 933578 (history)
Form: Version 2.0
Age: 43.0
Sex: Male
Location: New York
Vaccinated: 2021-01-08
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	012L20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: metform, atorvastin
Current Illness: Diabetes, Hypertension, Sleep Apnea, Obese

Preexisting Conditions: Hx smoking
Allergies: NKA
Diagnostic Lab Data:
CDC Split Type:

Write-up: Pronounced dead 1/9/2021 at 12:42. Received first dose of vaccine 1/8/2021

VAERS ID: 938097 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-12

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021017780

Write-up: died; This is a spontaneous report from a non-contactable consumer via a Pfizer-sponsored program. 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 medical history and concomitant medications were not reported. It was reported the patient was a doctor, died after the vaccine with no apparent disease. It was not reported if an autopsy was performed. No follow-up attempts are possible. Information about lot/batch number cannot be obtained.; Reported

Cause(s) of Death: Unknown cause of death

VAERS ID: 948181 (history)
Form: Version 2.0
Age: 89.0
Sex: Male
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Chest pain, Death, Heart rate irregular
SMQs:, Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Cardiac arrhythmia terms, nonspecific (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: ANORA, Albuterol, Flovent, Bumex, Carvediol; omeprazole
Current Illness:
Preexisting Conditions: COPD, CHF, renal; ASCVD; MI hx
Allergies: 0
Diagnostic Lab Data:
CDC Split Type:

Write-up: Death Chest pain; irreg heart rhythm; evening of vaccine; death on toilet on 1/13/21

VAERS ID: 955532 (history)
Form: Version 2.0
Age: 51.0
Sex: Female
Location: New Jersey
Vaccinated: 2021-01-12
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-01-19

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

Administered by: Senior Living **Purchased by:** ?
Symptoms: Cardioversion, Death, Posture abnormal, Pulse absent, Respiratory arrest, Resuscitation, Unresponsive to stimuli
SMQs:, Anaphylactic reaction (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Dystonia (broad), Acute central respiratory depression (narrow), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (broad), Respiratory failure (narrow), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-13
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Lantus SoloStar; Atorvastatin Calcium; Gabapentin; Potassium Chloride ER; Levothyroxine Sodium; Alogliptin Benzoate; Acetaminophen; Milk of Magnesia; Enema Rectal; Bisacodyl Rectal Suppository; Tramadol HCl; Ceftriaxone Sodium; Santyl Exter
Current Illness: Cellulitis of RLE, S/P wound debridement
Preexisting Conditions: HTN, Morbid obese, Diabetes, Hypothyroidism, HTN
Allergies: NKA
Diagnostic Lab Data: N/A
CDC Split Type:

Write-up: COVID 19 Vaccination administered by pharmacy staff. No adverse effect at the present time. Staff will continue to observe adverse reaction. Will continue to monitor. Patient at start of shift awake in the bed. Pt at 3am was on the commode leaned to the side. Patient body still warm to touch no pulse. Called for assistance Asap. Cpr started promptly. Cpr given patient on floor 911 arrived at the scene at 3:10am Cpr rotated Between Nursing and EMT on Scene. Cpr was given to patient for over 45 minutes. Patient was pronounced at the scene at 3:50am. Call placed to Pt family by supervisor on shift. MD to be notified. AT 3:00am, I was notified by the nurse that resident is unresponsive. Upon entering room, resident was sitting on the commode unresponsive with absent respiration and pulse. Resident lowered down on the floor with 4 person assist. CPR initiated, AED pads placed on chest with no shock indicated. 911 called and EMT and paramedics arrived around 3:10am. ACLS performed until code stopped and pronounced death at 3:48am. I called and notified family member of his demise and awaiting for family to call us back for funeral arrangements.

VAERS ID: 955879 (history)
Form: Version 2.0

Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-19

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034599

Write-up: expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar death events for 8 patients. This report is for 5th of 8 patients. A patient of unspecified age and gender received the first dose of 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 and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Sender''s Comments: The limited information provided in this report does not allow a full assessment of the case. The event death with unknown cause is assessed as related to the suspect drug per company guidance. This case will be reassessed when additional information, particularly the clinical course before death, complete medical history and concomitant medication and autopsy report, becomes available.,Linked Report(s) : US-PFIZER INC-2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: expired before receiving the second dose

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VAERS ID: 955880 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034603

Write-up: passed unexpectedly; This is a spontaneous report from a contactable nurse communicated to a Pfizer colleague. This nurse reported similar death events for 8 patients. This report is for 8th of 8 patients. A patient of unspecified age and gender 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 and concomitant medications were not reported. The patient passed unexpectedly on an unspecified date. The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Sender''s Comments: The limited information provided in this report does not allow a full assessment of the case. The event death with unknown cause is assessed as related to the suspect drug per company guidance. This case will be reassessed when additional information, particularly the clinical course before death, complete medical history and concomitant medication and autopsy report, becomes available. The case will be reassessed if 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.,Linked Report(s) : US-PFIZER INC-

2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: passed unexpectedly

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VAERS ID: 960426 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034595

Write-up: expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar death events for 8 patients. This report is for 1st of 8 patient. A patient of unspecified age and gender received first dose of 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 and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It was not reported if an autopsy was performed. Information about Lot/Batch number is requested.; Sender''s Comments: Current information is very limited for full assessment. The patient died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details especially death cause and autopsy results are needed for meaningful evaluation. 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) : US-PFIZER INC-2021034597 same drug, reporter and event but different patient;US-PFIZER INC-2021034598 same drug, reporter and event but different patient;US-PFIZER INC-2021034599 same drug, reporter and event but different patient;US-PFIZER INC-2021034600 same drug, reporter and event but different patient;US-PFIZER INC-2021034601 same drug, reporter and event but different patient;US-PFIZER INC-2021034603 same drug, reporter and event but different patient;US-PFIZER INC-2021034596 same drug, reporter and event but different patient.; Reported Cause(s) of Death: expired before receiving the second dose

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VAERS ID: 960427 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034596

Write-up: expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 2nd of 8

patients. A patient of unspecified age and gender received the first dose of 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 and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It was not reported if an autopsy was performed. Information about Lot/Batch number is requested.; Sender''s Comments: Current information is very limited for full assessment. The patient died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details especially death cause and autopsy results are needed for meaningful evaluation. 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.; Reported Cause(s) of Death: expired before receiving the second dose

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VAERS ID: 960428 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC2021034597

Write-up: expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 3rd of 8 patients. A patient of unspecified age and gender received the first dose of 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 and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It was not reported if an autopsy was performed. Information about Lot/Batch number is requested.; Sender"s Comments: Based on the reasonable temporal association, the Company cannot completely exclude the possible causality between the reported death and the administration of COVID 19 vaccine, bnt162b2. However, more information on the patient"s underlying medical condition, concomitant medications, patient"s age group, clinical course and relevant lab tests would be helpful for the Company to make a more meaningful causality 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 RA, IEC, as appropriate.; Reported Cause(s) of Death: expired before receiving the second dose

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VAERS ID: 960429 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:

Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034598

Write-up: expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 4th of 8 patient. A patient of unspecified age and gender received the first dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 22Dec2020 at single dose for covid-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information about Lot/Batch number has been requested.; Sender''s Comments: Current information is very limited for full assessment. The patient died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details especially death cause and autopsy results are needed for meaningful evaluation. 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) : US-PFIZER INC-2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: expired before receiving the second dose

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VAERS ID: 960430 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034600

Write-up: 7 residents expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 6th of 8 patients. A patient of unspecified age and gender received first dose 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''s medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Sender''s Comments: The event death is assessed as related to BNT162b2 vaccine and documented as such in the global safety database until sufficient information is available to allow an unrelated causality 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.,Linked Report(s) : US-PFIZER INC-2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: 7 residents expired before receiving the second dose

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VAERS ID: 960431 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 2020-12-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021034601

Write-up: expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar death events for 8 patients. This report is for 7th of 8 patient. A patient of unspecified age and gender received first dose of 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 and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It was not reported if an autopsy was performed. Information about Lot/Batch number is requested.; Sender''s Comments: Current information is very limited for full assessment. The patient died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details especially death cause and autopsy results are needed for meaningful evaluation. 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) : US-PFIZER INC-2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: expired before receiving the second dose

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VAERS ID: 962308 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Utah
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-21

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Dementia; Hospice care (on hospice, frail, but in good condition)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021045659

Write-up: died; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that an 83-year-old female patient (reporter mother) received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), via an unspecified route of administration on an unspecified date at single dose for covid-19 immunization. Medical history included hospice care and dementia. The patient''s concomitant medications were not reported. The patient died one day after getting vaccine. She was reportedly in good health the day before receiving vaccine. She was on hospice, frail, but in good condition and checked by a hospice nurse the day before which she reported her in good health considering. She was with dementia but stable in her health. The reporter read investigating 23 deaths of people receiving vaccine in similar conditions. The patient died on an unspecified date. It was not reported if an autopsy was performed.; Reported Cause(s) of Death: died

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VAERS ID: 970043 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 2020-12-10
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Diarrhoea
SMQs:, Pseudomembranous colitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Noninfectious diarrhoea (narrow)

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 2020-12-10
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Diarrhoea; Gastrointestinal disorder; Pseudomembranous colitis
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021019157

Write-up: Reported causes of death: Diarrhoea; This is a spontaneous report from a contactable healthcare professional via agency and a non-contactable consumer via a Pfizer-sponsored program Corporate (Pfizer) Social Media Platforms. An elderly patient of an unspecified age (also reported as were in their early to mid-60''s) and gender received BNT162B2 (Pfizer-BIONTECH Covid-19 Vaccine), via an unspecified route of administration on 10Dec2020 at a single dose for COVID-19 immunisation. Medical history included pseudomembranous colitis (broad), gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), and noninfectious diarrhoea (narrow) . The patient''s concomitant medications were not reported. The patient experienced diarrhoea on an unspecified date in 2020. It was reported that most of the deaths after the COVID-19 vaccine occurred within 24-48 hours after the shot. The write-ups that accompanied the reports furnished details about these sad fatalities, including the astonishing fact that some of the decreased had actually experienced and recovered from COVID-19 (raising questions about why they were vaccinated). It was also reported that the event was not life-threatening, did not result to a birth defect or permanent disability, did not require any office/ER/doctor visit, and did not require any hospitalization. The patient died on 10Dec2020. It was not reported if an autopsy was performed. The reported cause of death: diarrhoea. No follow up attempts are possible, information about the lot/batch number cannot be obtained.; Sender''s Comments: Based on the available information the event diarrhea resulting in death is attributed to patients preexisting medical conditions including pseudomembranous colitis, gastrointestinal nonspecific symptoms and therapeutic procedures, and noninfectious diarrhea. However, based on a close chronological association (same day) contributory role of BNT162B2 (Pfizer-BIONTECH Covid-19 Vaccine) to event exacerbation 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.; Reported Cause(s) of Death: Reported causes of death: Diarrhoea

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VAERS ID: 970162 (history)
Form: Version 2.0
Age:
Sex: Male

Location: Maryland
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021046017

Write-up: received the vaccine on Tuesday and was found dead at his kitchen table Wednesday afternoon; This is a spontaneous report from a contactable consumer. An 89 years old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unknown date, at single dose, for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient received the vaccine on Tuesday (unspecified date) and was found dead at his kitchen table on Wednesday afternoon (unspecified date). Cause of death was unknown. It was unknown if an autopsy was performed. Information about batch/lot number has been requested.; Reported Cause(s) of Death: received the vaccine on Tuesday and was found dead at his kitchen table Wednesday afternoon

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VAERS ID: 971562 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-01-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Immune thrombocytopenia
SMQs:, 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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021058222

Write-up: died; acute immune thrombocytopenia; This is a spontaneous report from two contactable consumers. A patient of unspecified age and gender received BNT162B2(lot number and expiration date not provided) via an unspecified route of administration on unspecified date at single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient died after receiving the covid vaccine on an unknown date. The patient developed acute immune thrombocytopenia on an unknown date. It was unknown if autopsy was performed. The cause of death was unknown. The outcome of the event "died" was fatal and of the event " acute immune thrombocytopenia" was unknown. The reporter wondered if a platelets blood problem may lead to death and if who have a blood platelets condition like essential thrombocytosis should not risk taking the vaccine. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Died

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VAERS ID: 978873 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-01-27

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, SARS-CoV-2 test positive
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 202012; Test Name: Covid-19; Test Result: Positive
CDC Split Type: USPFIZER INC2021068689

Write-up: died several hours after receiving a Covid-19 vaccine; 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''s medical history and concomitant medications were not reported. The patient died several hours after receiving a Covid-19 vaccine on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The person had tested positive for the virus (Covid-19) in late Dec2020. Information on the batch/lot number has been requested.; Reported Cause(s) of Death: died several hours after receiving a Covid-19 vaccine

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VAERS ID: 978876 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-01-27

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Autoimmune disorder, Death
SMQs:, Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021070550

Write-up: Autoimmune disease; This is a spontaneous report from a Pfizer-sponsored program from a contactable nurse. A male patient of an unspecified age received bnt162b2 (BNT162B2, Manufacturer Pfizer-BioNTech), via an unspecified route of administration, on an unspecified date, at single dose, for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. The patient experienced autoimmune disease on an unspecified date. The patient died on an unspecified date due to autoimmune disease. It was unknown if an autopsy was performed. The information on the lot/batch number has been requested.; Sender"s Comments: The information available in this report is limited and does not allow a medically meaningful assessment of the case. The company cannot completely exclude a causal relationship between the fatal autoimmune disease and suspect vaccine BNT162B2. Additional information regarding therapy duration, relevant medical history, underlying conditions, concomitant medications and detailed clinical course around the event onset will aid in comprehensive 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.; Reported Cause(s) of Death: Autoimmune disease

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VAERS ID: 991450 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-01-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Muscle spasms, Muscle tightness, Myalgia, Neck pain
SMQs:, Rhabdomyolysis/myopathy (broad), Dystonia (broad), Eosinophilic pneumonia (broad), Arthritis (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Lisinopril, furosemide, omeprazole, amlodipine, meclizine, loratadine
Current Illness:
Preexisting Conditions: Coronary artery disease (CAD); Hypertension (HTN); Renal Disease (e.g. CKD, HD, ESRF)
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: The patient developed left sided neck and trapezoid tightness and pain after receiving the moderna covid vaccine in her left shoulder. The injection site is non tender and does not show any erythema or tenderness. The tenderness is over the left trapezius and left lateral neck area. It also feel tight like a muscle spasm.

VAERS ID: 993822 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021075824

Write-up: he got both doses then a few days later he died; This is a spontaneous report from a contactable consumer reporting for a friend"s father. A male patient of an unspecified age received the second dose of bnt162b2 (BNT162B2) vaccine , via an unspecified route of administration on an unspecified date at single dose for Covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient received the first dose on an unknown date. The patient died few days after receiving the second dose of the vaccine on an unspecified date. It was not reported if an autopsy was performed. Information about lot/batch number has been requested.; Reported Cause(s) of Death: he got both doses then a few days later he died

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VAERS ID: 993823 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-02

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

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

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021077709

Write-up: died due to heart attack; This is a spontaneous report from a contactable consumer (reporting for her son-in-law) from the Pfizer-sponsored program Pfizer First Connect. A male patient of an unspecified age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on an unspecified date at a single dose for COVID-19 immunization. The patients medical history and concomitant medications were not reported. The patient died due to heart attack on an unspecified date. The patient died on an unspecified date. It was unknown if an autopsy was performed. Follow-up (28Jan2021): This follow-up is being submitted to notify that the lot/batch number is not available despite the follow-up attempts made. Follow-up attempts completed. No further information is expected.; Reported Cause(s) of Death: died due to heart attack

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VAERS ID: 993832 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Maine
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-02

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021081901

Write-up: At the end of the conversation, caller stated that recently saw in (place name) that someone passed away 3 hours after receiving the injection.; This is a spontaneous report from a contactable nurse. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on unspecified date at single dose for COVID-19 immunization. Medical history and concomitant medication were not reported. Caller, calling on behalf of her sister who has pseudocholinesterase imbalance, would like to know if the Covid vaccine has any contraindication or interaction with succinylcholine. Caller stated that her and her sister are scheduled to receive the first dose of the vaccine this weekend. Caller stated that her sister had a severe reaction to succinylcholine and did not wake up for four days. Caller stated that her sister has to wear a medical bracelet because of this condition. Caller reported that her sister"s son has the same severe reaction to succinylcholine. Caller also stated that when she was a director in the lab, she heard of a person passing away in the OR due to the same reaction with succinylcholine. At the end of the conversation, caller stated that recently saw in (state name) that someone passed away 3 hours after receiving the injection. It was unknow if autopsy was done. Information on the lot/Batch number has been requested.; Sender"s Comments: The information currently available does not allow a medically meaningful assessment for the event "passed away" with unknown cause of death. 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.; Reported Cause(s) of Death: At the end of the conversation, caller stated that recently saw in (place name) that someone passed away 3 hours after receiving the injection.

VAERS ID: 1000233 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-04

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021098743

Write-up: just died; This is a Spontaneous report from a Pfizer Sponsored Program from a contactable consumer. A male patient of an unspecified age 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''s medical history and concomitant medications were not reported. The patient just died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: just died

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VAERS ID: 1000624 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021064098

Write-up: Pneumonia; This is a spontaneous report from a Pfizer-sponsored program via a contactable consumer (patient). A male patient of unspecified age (Age: 63; Unit: Unknown) received the first dose and second dose of BNT162B2, via unspecified routes of administration on unspecified dates at single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. After receiving both vaccines, patient still got pneumonia and he had seen many elderly died from pneumonia even after receiving the vaccine. Outcome of the event was not resolved. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

VAERS ID: 1001713 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data: Test Name: temperature; Result Unstructured Data: 98
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Passed away; A spontaneous report was received from a consumer concerning a patient who received Moderna"s COVID-19 Vaccine (mRNA-1273) and passed away. The patient"s medical history was not provided. No relevant concomitant medications were reported. On an unknown date, approximately 2 hours prior to the onset of the event, the patient received a dose of mRNA-1273 intramuscularly in the for prophylaxis of COVID-19 infection. They were not feeling sick or experiencing any adverse events. Vital signs included temperature 98 degrees Fahrenheit. Approximately two hours after receiving the vaccine, the patient passed away. No treatment information was provided. Action taken with the drug in response to the event was not applicable. The patient died on an undisclosed date. The cause of death was not provided. Plans for an autopsy were not provided.; Reporter"s Comments: This case concerns a patient of unknown age and gender. The medical history and concomitant medication is not provided. The patient experienced Death. The event occurred approximately two hours after receiving their first of two planned doses of mRNA-1273 (Lot unknown). Very limited information regarding this event has been provided at this time. Based on temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. The benefit-risk relationship of Moderna"s COVID-19 vaccine is not affected by this report.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1004645 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-05

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021096325

Write-up: Pfizer vaccine caused the death; This is a spontaneous report from a non-contactable consumer. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot number and expiration date were not reported), 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. The reporter claimed that Pfizer vaccine caused the death of the doctor (patient). No further details reported. No follow-up attempts are possible. Information about lot and batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: Pfizer vaccine caused the death

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VAERS ID: 1010899 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021102020

Write-up: On the two people who died, one in (State name) and one in (State name); This is a spontaneous report from a contactable other HCP. This other HCP reported similar events for 2 patients. This is 1 of 2 report. An unknown age male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecific date at single dose for covid-19 immunization. Medical history and concomitant drug were not reported. It was reported that patient was died. Cause of death unknown. Outcome of the event was fatal. Information on the lot/batch number has been requested.; Sender''s Comments: Death with unknown cause is considered related to BNT162B2 for reporting purpose. Information is very limited. Case will be reassessed once receiving additional information, including cause of death. 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) : US-PFIZER INC-2021102049 same reporter/drug/event, different patient; Reported Cause(s) of Death: On the two people who died, one in (State name) and one in (State name)

VAERS ID: 1010900 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021102049

Write-up: patient died; This is a spontaneous report from a contactable Other Healthcare Professional (HCP). This Other HCP reported similar event for 2 patients. This is 2nd of 2 reports. A male patient of an unspecified age 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"s medical history and concomitant medications were not reported. It was reported that patient died on an unspecified date. Cause of death unknown. Outcome of the event was fatal. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Sender"s Comments: The causal relationship between BNT162B2 and the event death cannot be excluded as 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.,Linked Report(s) : US-PFIZER INC-2021102020 same reporter/drug/event, different patient; Reported Cause(s) of Death: patient died

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VAERS ID: 1013095 (history)
Form: Version 2.0
Age: 88.0
Sex: Male
Location: Hawaii
Vaccinated: 2021-02-03
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	LA / SYR

Administered by: Senior Living **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 2021-02-04
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: See list below. Flomax 0.4 mg 1 Cap P.O. QD Tylenol 500mg 2 tabs PO PRN Carbidopa/levodopa 25-100mg PO 3 tab PO TID CoQ10 1 cap PO QD B12 1 tab PO QD Omega-3 Fatty Acids-Vitamin E 1,000mg 1 Cap PO QD Timolol 0.25% 1 gtt OU BID Finasterid
Current Illness: Unknown.
Preexisting Conditions: See list below. Glaucoma, Hyperlipidemia, Inguinal Hernia, Hx-Gi bleed, Mandibular salivary gland removal- 2012, Hx-TIA, Hypertension, Lumbar disc disease, Parkinsonism, Pancreatitis history
Allergies: No known drug allergies
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient was technically a resident of retirement community, however, he chose to live independently at home at his current home. Only his spouse actually resides at retirement community. Patient received the second dose of the Moderna vaccine via pharmacy vaccination clinic at retirement community on 2/3/2021. He was found deceased in his home by a certified nursing assistant on 2/4/2021 at approximately 10:00am.

VAERS ID: 1015630 (history)
Form: Version 2.0
Age: 56.0
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-09

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died of a heart attack; A spontaneous report was received from a consumer concerning a 56-year-old male patient who received Moderna"s COVID-19 Vaccine (mRNA-1273) and died due to a heart attack. The patient"s medical history was not provided. Products known to have been used by the patient, within two weeks prior to the event, included additional medications. On an unknown date, the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. - On an unknown date, the reporter stated the patient received Moderna"s COVID-19 vaccine and sometime after he was found dead by his bedside around 02:30 am. The patient died on an unknown date. The cause of death was due to a heart attack. Plans for autopsy were not provided.; Reporter"s Comments: This case concerns a 56 year old, male patient, who experienced fatal unexpected event of myocardial infarction. The event occurred on an unspecified date after mRNA-1273 (Lot# Unknown). Treatment details not provided. Very limited information regarding this event has been provided at this time. Based on temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Additional information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1016605 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-09

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Sepsis
SMQs:, Sepsis (narrow), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Sepsis; A spontaneous report was received from a consumer post , concerning an approximately 55-year-old, male physician who received Moderna"s COVID-19 vaccine (mRNA-1273) and developed sepsis, resulting in death. There was no medical history provided. There were no concomitant medications provided. On an unknown date (Thursday), the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. According to the post, two days after vaccine administration, the patient died of sepsis on Saturday. Action taken with mRNA-1273 in response to the event was not applicable. The event, sepsis, was considered fatal. The patient"s date of death was not provided. The cause of death was reported as sepsis.; Reporter"s Comments: This case concerns a 55-year-old, male subject, who experienced a serious unexpected event of Sepsis. Sepsis occurred after first dose of mRNA-1273 vaccine administration. On an unknown date, two days after vaccine administration, the patient died of sepsis. Treatment for the event was not provided. The patient"s medical history was not provided. The patient is a physician. Concomitant product use was not reported. Very limited information regarding this event has been provided at this time and no definite diagnosis or autopsy report have been provided. Based on the current available information and temporal association between the use of the product and the start date of the event of Sepsis, a causal relationship cannot be excluded.; Reported Cause(s) of Death: Sepsis

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VAERS ID: 1017128 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021130110

Write-up: Passed away; This is a spontaneous report from a Pfizer-sponsored program Corporate (Pfizer) Social Media Platforms. This is a spontaneous report from a contactable consumer reporting for friend"s mother. A 50-years-old female patient received the second dose of bnt162b2 (BNT162B2) vaccine , via an unspecified route of administration on an unspecified date at single dose for Covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient received the first dose of BNT162B2 vaccine on an unknown date. The patient passed away on an unspecified date. The patient was a healthy woman, who just got her 2nd dose of the vaccine a couple of days before. The patient died in her sleep. Doctor labeled her death as "natural causes". It was not reported if an autopsy was performed. No follow-up attempts are Possible. Information on lot/batch cannot be obtained.; Reported Cause(s) of Death: Passed away

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VAERS ID: 1026021 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Maryland
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	- / -
PPV: PNEUMO (NO BRAND NAME) / UNKNOWN MANUFACTURER	- / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Inappropriate schedule of product administration
SMQs:, Medication errors (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021120769

Write-up: a male patient received the pneumonia shot 12 days after the first dose of the vaccine; he had died; This is a spontaneous report from a contactable consumer or other non hcp. A 76 years old male patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date at single dose for COVID 19 immunisation and received pneumococcal 13-val conj vac (dipht crm197 protein) (PNEUMOCOCCAL 13-VAL CONJ VAC (DIPHT CRM197 PROTEIN)), via an unspecified route of administration on an unspecified date 12 days after bnt162b2 at single dose for immunisation. The patient medical history and concomitant medications were not reported. The patient died on an unspecified date. It was not reported if an autopsy was performed. Pfizer is Marketing Authorization Holder of pneumococcal 13-val conj vac (dipht crm197 protein) in the reporter''s country. This may be a duplicate report in situations where another Marketing Authorization Holder of pneumococcal 13-val conj vac (dipht crm197 protein) has submitted the same report to the regulatory authorities. Information on lot and batch number has been requested. Follow-up: (08Feb2021): Lot/batch number is not available despite the follow-up attempts made. Follow-up attempts are completed. No further information is expected.; Reported Cause(s) of Death: a male patient received the pneumonia shot 12 days after the first dose of the vaccine; he had died

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VAERS ID: 1030011 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: ELIQUIS
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021149056

Write-up: taking Eliquis who died after receiving the Pfizer-BioNtech Covid-l9; This is a spontaneous report from a contactable consumer based on information received by Pfizer from Bristol-Myers Squibb (manufacturer control number US-BRISTOL-MYERS SQUIBB COMPANY-BMS-2021-014171), license party for apixaban (ELIQUIS). This spontaneous case was reported by a non-health professional and describes the occurrence of DEATH (taking Eliquis who died after receiving the Pfizer-BioNtech Covid-l9) in patient of an unknown age and gender who received apixaban (Eliquis) for an unknown indication. CO-SUSPECT PRODUCTS included Covid-19 Vaccine. On an unknown date, the patient started Eliquis (unknown route) and Covid-19 Vaccine (unknown route). DEATH occurred on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. The doctor died after taking Eliquis with Covid-19 Vaccine. For Eliquis(Unknown), the reporter did not provide any causality assessments. This case was linked to US-BRISTOL-MYERS SQUIBB COMPANY-BMS-2021-012621 (Linked Report).; Sender''s Comments: BMS Medical Evaluation Comment: This patient died after receiving apixaban therapy. Patient also received COVID-19 vaccine. Based on the limited information available regarding the cause of death and autopsy details, it cannot be ascertained with the reasonable possibility that the apixaban could have caused the event.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1030025 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Oklahoma
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No adverse event history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Death; A spontaneous report was received from a consumer concerning a male patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and the patient was died. The patient"s medical history was not provided. Concomitant product use was not provided. On an unknown date, the patient received his first dose of mRNA-1273 (Lot number unknown) for prophylaxis of COVID-19 infection. On an unknown date, the patient was died. Treatment of this event was not provided. The patient was died. The cause of death was not provided. Autopsy details were not provided.; Reporter"s Comments: This case concerns a male patient (unknown age), who experienced event of death (cause unknown). The event occurred on an unknown date after the first and last dose of mRNA-1273 vaccine administration. Autopsy and cause of death were not reported. Based on the current available limited information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded and the events are assessed as possibly related.; Reported Cause(s) of Death: Death

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VAERS ID: 1030132 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebral haemorrhage, Death, Platelet count decreased
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Haemorrhagic central nervous

system vascular conditions (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: platelet count; Result Unstructured Data: Test Result:low platelet
CDC Split Type: USPFIZER INC2021135267

Write-up: a doctor died of low platelet and brain bleed 16 days after the vaccine; a doctor died of low platelet and brain bleed 16 days after the vaccine; The 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 medical history was not reported. The patient"s concomitant medications were not reported. The patient (doctor) died of low platelet and brain bleed 16 days after the vaccine. It was not reported if an autopsy was performed. Information about lot/batch number has been requested.; Reported Cause(s) of Death: a doctor died of low platelet and brain bleed 16 days after the vaccine; a doctor died of low platelet and brain bleed 16 days after the vaccine

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VAERS ID: 1030852 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; A spontaneous report was received from a reporter concerning a unknown patient who received Moderna"s COVID-19 Vaccine (mRNA-1273) and passed away. The patient"s medical history was not provided. No relevant concomitant medications were reported. On an unknown date, the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. On an unknown date, the patient passed away. No treatment information was provided. Action taken with RNA-1273 in response to the event was not applicable. The patient died on unknown date. The cause of death was reported as unknown. Autopsy details were unknown.; Reporter"s Comments: This case concerns a patient of unknown age and gender. The patient"s medical history was not provided. The fatal, unexpected event of death occurred on an unknown date after the administration of the first dose of mRNA-1273 on an unknown date. The cause of death was reported as unknown. Autopsy details were unknown. Very limited information regarding this event has been provided at this time. Based on temporal association between the use of the product and the event, a causal relationship cannot be excluded. Additional information regarding the autopsy report, date of the mRNA administration, day of death, medical information and details of concomitant product are all required for further assessment of causality.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1033466 (history)
Form: Version 2.0
Age: 70.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest, Death, Resuscitation
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Asthma/COPD, Diabetes mellitus (DM), Hypertension (HTN), Renal Disease (e.g. CKD, HD, ESRF), A-fib, pulmonary hypertension, chronic loculated empsyema, chronic aspiration pneumonitis, pacemaker, chronic anticoagulation
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: 70 yo man with multiple severe medical comorbidities received his first dose of Moderna COVID-19 vaccination without incident. 8.5 hours later, he was noted by his family to be in his usual state of health. 9.5 hours after the vaccination, he was found down by his family in cardiac arrest. Resuscitation attempts were not successful, and the patient died.

VAERS ID: 1033472 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Resident coded and expired; A spontaneous report was received from a consumer concerning a patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and coded and expired. The patient"s medical history was not provided. No concomitant product use was reported. On an undisclosed date, the patient received their first of two planned doses of mRNA-1273 (Lot number: not provided) for prophylaxis of COVID-19 infection. On undisclosed date, the patient coded and expired. Treatment information was not provided. Action taken with mRNA-1273 in response to the events was not applicable. The patient died on an undisclosed date. The cause of death was not provided. Plans for an autopsy were not provided.; Reporter"s Comments: MEDICAL COMMENT: MOD 2021 009822 DEATH NOS This case concerns a patient of unknown age and gender who received their first of two planned doses of mRNA-1273 (Lot number: not provided) for prophylaxis of COVID-19 infection and had died. Very limited information regarding this event has been provided at this time to make a proper medical assessment, therefore, the causality is unlikely related to the vaccine in this case of death not otherwise specified. No contact information provided. Follow up is not expected.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1034985 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Kentucky
Vaccinated: 2021-01-04
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-17

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	039K20A / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: 4 Nurshing home patients Died; A Spontaneous report was received from a pharmacist concerning 4 nursing home patients of unspecified age and gender who received Moderna"s COVID-19 vaccine (mRNA-1273) and died. The patients" medical histories were not provided. No relevant concomitant medications were reported. On unspecified dates, 4 nursing home patients received their first of two planned doses of mRNA-1273 (Lot # 039K20A) for prophylaxis of COVID-19 infection. A pharmacist reported that they just learned 4 nursing home patients died after the first dose of the Modern vaccine. The patients were buried, and no autopsies were conducted. The pharmacist suspected latent Covid-19 on the patients and that the vaccine precipitated this outcome. No treatment information was provided. Action taken with the second dose of mRNA-1273 in response to the event was not applicable. The event 4 nursing home patients died was fatal.; Reporter"s Comments: This case concerns 4 nursing home patients of unspecified age and gender who received their first dose of Moderna"s COVID-19 vaccine (mRNA-1273)Lot # 039K20A) and died. Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: died

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VAERS ID: 1035505 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-17

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Passed away with an hour to hour and 1/2 of receiving vaccine; A spontaneous report was received from a consumer concerning a patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and passed away with an hour to hour and 1/2 of receiving vaccine. The patient"s medical history, as provided by the reporter, included COVID-19. No relevant concomitant medications were reported. On unknown date, prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. On an unknown date, the patient passed away within an hour and a 1/2 of receiving the vaccine. Per the nursing home staff, they did not expect the patient to make it many more days. The patient was unresponsive in the room when the shot was given. The patient was 14+ days post COVID. No treatment information was provided. Action taken with mRNA-1273 in response to the event was not applicable. The patient died on an unknown date. The cause of death was unknown.; Reporter"s Comments: This case concerns a patient, who experienced event of death (unknown cause). The event occurred an hour to hour and 1/2 after the first and last dose of mRNA-1273 vaccine administration. Based on the current limited available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded and the event is assessed as possibly related. However, Per the nursing home staff, the patient was 14+ days post COVID and they did not expect the patient to make it many more days. The patient was unresponsive in the room when the shot was given.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1035545 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-17

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021139379

Write-up: Death; This is a spontaneous report from a contactable consumer. A 42-year-old male patient received Covid 19 Vaccine (UNSPECIFIED TRADE NAME), via an unspecified route of administration from an unspecified date to an unspecified date at single dose for covid-19 immunization.The patient medical history was not reported. The patient''s concomitant medications were not reported. The reporter was calling about the Covid 19 Vaccine. Reporter stated that she gets information every day from the nursing home. Reporter stated that she will provide the name of the Nursing home, she just forgot to put it down. The patient, within 2 days dead from it, he was perfectly healthy. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on Lot/batch number has been requested.; Reported Cause(s) of Death: Death

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VAERS ID: 1035549 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-17

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Blood bilirubin, Blood osmolarity, Blood pressure measurement, Blood urea, Monocyte count, Platelet count, Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Systemic lupus erythematosus (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: blood bilirubin; Result Unstructured Data: Test Result:1.50; Test Name: osmolality; Result Unstructured Data: Test Result:297; Test Name: blood pressure; Result Unstructured Data: Test Result:159/106; Test Name: Blood urea (BUN); Result Unstructured Data: Test Result:22.3; Test Name: monocytes; Test Result: 12.1 %; Test Name: platelet count; Result Unstructured Data: Test Result:Thrombocytopenia; Test Date: 20201221; Test Name: platelet count; Result Unstructured Data: Test Result:platelet count decreased again to 0
CDC Split Type: USPFIZER INC2021144615

Write-up: Thrombocytopenia; This is a spontaneous report from a contactable consumer (patient"s wife). A male patient of unspecified age 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 medical history and concomitant medications were not reported. The patient experienced thrombocytopenia on an unspecified date, the seriousness of the event reported as death. The patient died two weeks after receiving a COVID-19 vaccine. Patient"s wife said that he died from a condition known as thrombocytopenia, marked by a shortage of the blood platelets that help stop bleeding, after he received BNT162B2. The patient underwent lab tests and procedures which included blood bilirubin: 1.50, blood osmolarity: 297, blood pressure measurement: 159/106, blood urea: 22.3, monocyte count: 12.1 %, platelet count: thrombocytopenia, platelet count: platelet count decreased again to 0 on 21Dec2020. The patient died on an unspecified date. It was unknown if an autopsy was performed. The outcome of the events was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Thrombocytopenia

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VAERS ID: 1037720 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Completed suicide

SMQs:, Suicide/self-injury (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Suicide; A spontaneous report was received from a consumer concerning a patient, of unknown age/gender, who received Moderna"s COVID-19 vaccine and experienced suicide. The patient"s medical history was not provided. Concomitant product use was not provided by the reporter. On unknown date, the patient received mRNA-1273 (Lot number: unknown) intramuscularly for prophylaxis of COVID-19 infection. On an unknown date the patient experienced suicide. Treatment information was not provided. Action taken with the mRNA-1273 in response to the event was not reported. The patient died on unknown date. The cause of death was reported as suicide. Plans for an autopsy were not provided.; Reporter"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Suicide

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VAERS ID: 1037867 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021153879

Write-up: Caller"s mother received both doses of the Pfizer covid vaccine; within a month of receiving the vaccine, the caller"s mother died.; This is a spontaneous report from a contactable consumer reported for the mother. A female patient of unknown age received both doses of vaccine BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), on unknown dates at single dose for COVID-19 immunization. Medical history and Concomitant medications were not reported. Within a month of receiving the vaccine, the patient died. It was unknown if an autopsy was performed. Information on the Lot/Batch number has been requested ; Reported Cause(s) of Death: within a month of receiving the vaccine, the caller"s mother died

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VAERS ID: 1038234 (history)
Form: Version 2.0
Age: 84.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-01-14
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-18

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

Administered by: Senior Living **Purchased by:** ?
Symptoms: Condition aggravated, Death, Dementia, Mental impairment, Pyrexia, Urinary tract infection, Wrong product administered
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Medication errors (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 2021-02-13
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Cephalexin Spiriva, Nitro, Probiotic, Albuterol, Vitamin D3, Multivitamin Aspirin, Glucosamine, Metamucil, Tylenol, Cranberry, Claritin, Quetiapine
Current Illness: Dementia with mental status change Urinary tract infection
Preexisting Conditions: Dementia Hypertension, Chronic UTI Hyperlipidemia, h/o Alcohol Abuse in Remission
Allergies: Sulfa, Aricept
Diagnostic Lab Data:
CDC Split Type:

Write-up: First dose Pfizer given at assisted living on 1/14/21 she was subsequently admitted to hospital and got her second dose but Moderna was given instead of Pfizer on 2/8/21. She had been improving but in the days following the Moderna vaccine she developed fever and then her mental status declined. She was discharged back to assisted living. Suspected UTI, and moderate dementia, placed on hospice (2/12/21). Died on 2/13/21.

VAERS ID: 1039952 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Off label use
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021155151

Write-up: Two report of death related to the vaccine were reported in a young premature baby and 25-year-old young person who dies 15 minutes after receiving the vaccine.; a young premature baby received BNT162B2; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported similar event for 2 patients. This is 1st of 2 reports. A patient of unspecified age and gender (reported as a young premature baby) received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The relevant medical history and concomitant medications were not reported. The consumer stated his main concern was even though the vaccine had been authorized. The vaccine was only given vaccine on the side of the road. People were dying from the vaccine more than from the virus. He gave out some percentage of people that could die from the virus vs from the vaccine. The percentage of death rate was 0.00045% from the virus. The percentage of death from the vaccine was 6.6%. There were many people reporting side effects such as bell''s palsy, whole body convulsion. Two report of death related to the vaccine were reported in a young premature baby and 25-year-old young person who died 15 minutes after receiving the vaccine. He said it''s sad these were front line worker who were facing all these side effects. The outcome of the event was fatal. Information on the lot/batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021157531 same reporter, drug, event, different patient; Reported Cause(s) of Death: Two report of death related to the vaccine were reported in a young premature baby and 25-year-old young person who dies 15 minutes after receiving the vaccine.

VAERS ID: 1039954 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021157531

Write-up: Two report of death related to the vaccine were reported in a young premature baby and 25-year-old young person who dies 15 minutes after receiving the vaccine.; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported similar event for 2 patients. This is 2nd of 2 reports. A 25-year-old patient of unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The relevant medical history and concomitant medications were not reported. The consumer stated his main concern was even though the vaccine had been authorized. The vaccine was only given vaccine on the side of the road. People were dying from the vaccine more than from the virus. He gave out some percentage of people that could die from the virus vs from the vaccine. The percentage of death rate was 0.00045% from the virus. The percentage of death from the vaccine was 6.6%. There were many people reporting side effects such as bell''s palsy, whole body convulsion. Two report of death related to the vaccine were reported in a young premature baby and 25-year-old young person who died 15 minutes after receiving the vaccine. He said it''s sad these were front line worker who were facing all these side effects. The outcome of the event was fatal. Information on the lot/batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021155151 same reporter, drug, event, different patient; Reported Cause(s) of Death: Two report of death related to the vaccine were reported in a young premature baby and 25-year-old young person who dies 15 minutes after receiving the vaccine.

VAERS ID: 1048665 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Internal haemorrhage

SMQs:, Haemorrhage terms (excl laboratory terms) (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; Internal bleeding and died; A spontaneous report was received from a nurse concerning a male patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and experienced internal bleeding and died. The patient"s medical history was not provided. No concomitant product use was reported. On unknown date, the patient received their first of two planned doses of mRNA-1273 (Lot number: not provided) intramuscularly for prophylaxis of COVID-19 infection. On unknown date, a nurse reported she read about a doctor who received the COVID-19 vaccine and died 3 days later. She stated she read that the doctor had internal bleeding. It started with bleeding in his hands and feet, then went to his brain and died 3 days later. Treatment information was not provided. Action taken with mRNA-1273 in response to the events was not provided. The outcome of the events, internal bleeding and died, was considered fatal.; Reporter"s Comments: Very limited information regarding the events has been provided at this time and is insufficient for causality assessment. Further information has been requested.; Reported Cause(s) of Death: Internal bleeding

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VAERS ID: 1048672 (history)
Form: Version 2.0
Age: 89.0
Sex: Male
Location: Unknown
Vaccinated: 2020-12-29
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Death
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: No adverse event, Continue: [UNK], Comment: No reported medical history.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Pass away; Got COVID; A spontaneous report was received on from a consumer, concerning an 87-year-old male patient, who received Moderna''s COVID-19 (mRNA 1273) vaccine and experienced pass away (death) and COVID (COVID-19). The patient''s medical history was not provided. No relevant concomitant medications were reported. On 29 Dec 2020, the patient received their first of two planned doses of mRNA-1273 injection for the prophylaxis of COVID-19 infection. Two days after mRNA-1273 injection, the patient''s daughter was diagnosed with COVID-19 infection. The daughter states that her mother also got infected with COVID. On an unknown date, the patient got COVID. It was reported that he passed away. Treatment for the event was not provided. The cause of death was not provided. The plans for an autopsy were not provided. .; Reporter''s Comments: Although the onset date of COVID-19 and the fatal outcome is lacking, Based on the natural history of COVID-19, the reported event, it is assessed as unlikely related with mRNA-1273. The actual cause of death cannot be ascertained but assessed as not related and most likely due to the COVID-19 infection.

VAERS ID: 1048686 (history)
Form: Version 2.0
Age:
Sex: Female
Location: California
Vaccinated: 2021-01-01
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: SARS-CoV-2 antibody test, Sepsis

SMQs:, Sepsis (narrow), Opportunistic infections (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Depression; Schizophrenia
Allergies:
Diagnostic Lab Data: Test Date: 202011; Test Name: covid19; Test Result: Positive
CDC Split Type: USPFIZER INC2021166424

Write-up: died just 10 days after being given the vaccine/ put sepsis on her medical records; This is a spontaneous report from a contactable consumer report for Aunt. A 59-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration in Jan2021 at single dose for COVID-19 immunization. Medical history included schizophrenia and depression. The patient''s concomitant medications were not reported. Consumer''s aunt (patient) was housed in a facility. She was being treated for schizophrenia and depression. This was one of the facilities that chose to house Covid patients during the pandemic. Many of the patients here contracted covid19 during this time and they had to do a full facility lockdown and quarantine. The patient tested positive in Nov2020 and was quarantined to her room for 10 days. 3 weeks ago (in Jan2021), she received the Pfizer vaccine. Consumer''s family was not given notice of this and we are sad to report that she died just 10 days after being given the vaccine. They put sepsis on her medical records and have not connected this to the vaccine. Consumer stated aunt was just 59 yrs old, though she was being treated for her mental illness, she was physically healthy. Consumer''s family gravely concerned that this facility neglected her health by administering the vaccine without considering possible reactions from the medication she was taking, or the fact she had Covid just months prior. Patient died on an unspecified date. it was unknown if an autopsy performed. Information on Lot/Batch number has been requested.; Reported Cause(s) of Death: died just 10 days after being given the vaccine/ put sepsis on her medical records

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VAERS ID: 1049284 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Unevaluable event (underlining health issue)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; A spontaneous report was received from a consumer concerning a 58-year-old, female patient who received Moderna''s COVID-19 vaccine (mRNA-1273) and patient was died. The patient''s medical history was reported as underlying health issue. Concomitant product use was not provided by the reporter. On an unknown date, prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (lot/batch: if reported or unknown) intramuscularly for prophylaxis of COVID-19 infection. On an unknown date, patient with underlying health issue died after getting the Moderna COVID-19 vaccine. Action taken with mRNA-1273 in response to the events was not applicable. The patient died on an unknown date. The cause of death was not provided. Plans for an autopsy were not provided.; Reporter''s Comments: This case concerns a 58-year-old female who died on unknown date after first dose of mRNA-1273, lot # unknown. Very limited information regarding this event has been provided at this time, therefore it is difficult to assess a cause and effect relationship. No further information will be available. Of note, patient''s medical history included unknown underlying health issues.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1049773 (history)
Form: Version 2.0
Age: 51.0
Sex: Female
Location: Georgia
Vaccinated: 2021-02-12
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-23

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	015M20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Death, Pulmonary embolism
SMQs:, Embolic and thrombotic events, venous (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: none listed on consent form
Diagnostic Lab Data: unknown
CDC Split Type:

Write-up: Patient died on 02/20/2021. Cause of death was pulmonary embolism.

VAERS ID: 1051451 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021179726

Write-up: died 15 min later; This is a spontaneous report from a Pfizer Sponsored Program. A non-contactable consumer reported that a female patient (mother) of an unknown age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) at single dose via an unknown route on an unknown date for Covid-19 immunization. Medical history and concomitant drug were not provided. The reporter stated her mother took the vaccine and died 15 min later. No follow-up attempts are possible. Information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: died 15 min later

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VAERS ID: 1059622 (history)
Form: Version 2.0
Age: 85.0
Sex: Male
Location: Massachusetts
Vaccinated: 2021-02-02
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL3247 / 2	AR / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Death, Metabolic encephalopathy, Resuscitation, Unresponsive to stimuli, Urinary tract infection, Vomiting
SMQs:, Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (narrow), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-08
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Lipitor Renvela caps, Pantoprazole, Humalog, Midodrine, Neptro caps, Metoprolol, Levemir, Levothyroxine
Current Illness: UTI & metabolic encephalopathy
Preexisting Conditions: Chronic Kidney Di Stage 5 with Renal Dialysis Diabetes Type 2, CHF, H/o Bladder CA, HTN
Allergies: Ace Inhibitors
Diagnostic Lab Data:
CDC Split Type:

Write-up: Resident rec"d COVID vaccine #2 on 02/02/2021 and was hospitalized on 02/03/2021. Diagnosed with UTI & Metabolic Encephalopathy. He was re-admitted to facility on 02/05/2021. On 02/08/2021 resident was found to be unresponsive with small amount of tan emesis in mouth and on bed. CPR initiated and resident was transferred to ER. ER MD notified facility that resident had died.

VAERS ID: 1056659 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-26

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac disorder
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021186885

Write-up: heart issue; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age 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 medical history and concomitant medications were not reported. The patient passed away after taking the vaccine. He was healthy but developed heart issue after taking vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed.; Reported Cause(s) of Death: heart issue

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VAERS ID: 1056669 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Nevada
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-26

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021203949

Write-up: She knows one person did die; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received BNT162B2, via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The reporter

mentioned that "she knows one person did die" on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: She knows one person did die

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VAERS ID: 1057547 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-26

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: 86 year old manager received the two doses of the Moderna vaccine and died; A spontaneous report was received from a consumer, concerning an 86-year-old, male patient who received Moderna''s COVID-19 vaccine (mRNA-1273) and passed away. The patient''s medical history was not provided. No relevant concomitant medications were reported. On an unspecified date, the patient received their doses of mRNA-1273 (Lot number: unknown) through an unknown route in an unknown arm for prophylaxis of COVID-19 infection. On an unspecified date, it was reported that the patient passed away after receiving both doses. The cause of death was unknown. It was unknown if an autopsy was performed. The patient received both scheduled doses of mRNA-1273 prior to the event; therefore, action taken with the drug in response to the events is not applicable. The outcome of event "86 year old manager received the two doses of the Moderna vaccine and died" was fatal.; Reporter''s

Comments: Very limited information regarding this event has been provided at this time. No autopsy report provided. Further information has been requested.; Reported Cause(s) of Death: 86 year old manager received the two doses of the Moderna vaccine and died

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VAERS ID: 1062961 (history)
Form: Version 2.0
Age: 90.0
Sex: Male
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-02-26

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: veltassa
Current Illness:
Preexisting Conditions: hypothyroidism, benign prostatic hypertrophy, hyperkalemia, stage 5 CKD on dialysis
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Called patient to refill medication and spoke with daughter. She stated that her father had passed away last week.

VAERS ID: 1061910 (history)
Form: Version 2.0
Age: 65.0
Sex: Male

Location: Pennsylvania
Vaccinated: 2021-02-11
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	030M20A / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-01
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Passed away; A spontaneous report was received from a Pharmacist concerning a 65 years-old, male patient who passed away/MedDRA PT: death, days after receiving the second dose of the Moderna COVID-19 vaccine. The patient"s medical history was not provided. Concomitant product use was not provided by the reporter. The patient received their first of two planned doses of mRNA-1273 (Lot #012L20A) on 12 Jan 2021. On 11 Feb 2021, the patient received their second of two planned doses of mRNA-1273 (Lot # 030M20A) (route of administration and injection site not provided) for prophylaxis of COVID-19 infection. On an unknown date, days after receiving the second dose of the Moderna COVID-19 vaccine, the patient passed away. The patient did not come to the hospital; therefore the Pharmacist had very little detail of the situation but believed it was due to aspiration based on report received from the patient"s boss. The Pharmacist reported there was no report of any issues from the first vaccine. The patient was found at home by spouse and had yellow stuff on face and chest. Action taken with mRNA-1273 was not applicable as the patient deceased. The patient died on an unknown date. The cause of death was reported as unknown. Plans for autopsy were not provided.; Reporter"s Comments: Very limited information regarding the event of death has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Death days after receiving the second dose of the Moderna Covid-19 Vaccine

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VAERS ID: 1065394 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-02

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; A spontaneous report was received from a consumer on a social media, concerning a 58-years-old female patient, unknown race and ethnicity, who was administered Moderna"s COVID-19 vaccine (mRNA-1273), and died. The patient"s medical history was not provided. No relevant concomitant medications were reported. On an unknown date, prior to the onset of the event, the patient received dose of mRNA-1273 (Lot number: unknown), for the prophylaxis of COVID-19 infection. On 17-Feb-2021, social media interaction was posted concerning a death of a patient on an unknown date after receiving Moderna vaccine. No treatment information was provided. Action taken with mRNA-1273 in response to the event was not applicable. The patient died on an unknown date. The cause of death was not provided. Plans for autopsy were not provided.; Reporter"s Comments: Very limited information regarding the event has been provided at this time and is insufficient for causality assessment. The cause of death was not provided. Plans for autopsy were not provided.; Sender"s Comments: US-MODERNATX, INC.-MOD-2021-018302:Same Reporter; Reported Cause(s) of Death: Unknown cause of death

---

VAERS ID: 1065765 (history)
Form: Version 2.0

Age: 78.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-18
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-02

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, SARS-CoV-2 test negative
SMQs:, COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: No known side effects; postmortem Covid-19 test negative Date of death: 02/16/2021

VAERS ID: 1066291 (history)
Form: Version 2.0
Age: 80.0
Sex: Female
Location: Louisiana
Vaccinated: 2021-01-14
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	011J204 / 1	RA / IM

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

Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-30
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: nitroglycerin, asa,crestor,lantus,gabapentin,clopidogrel isosorbide, metformin,metoprolol, furosemide, ramipril, pantoprazole, albuterol, ondansetron, methocarbamol, norco, diclofenac Patient was given influenza vaccine by an outside provid
Current Illness: none
Preexisting Conditions: Type 2 diabetes, heart failure, hyperlipidemia, claudication, left hip and knee pain
Allergies: Aspirin, pneumococccal vaccine
Diagnostic Lab Data: N/A
CDC Split Type:

Write-up: Patient administered vaccine on 1/14/21 with no reactions noted after 15 minutes. Patient seen for wellness visit on 1/20/21 with no complaints regarding vaccine. Patient died on 1/30/21

VAERS ID: 1066307 (history)
Form: Version 2.0
Age: 95.0
Sex: Female
Location: Louisiana
Vaccinated: 2021-01-27
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9265 / 1	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-11
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknown
Preexisting Conditions: unknown
Allergies: unknown
Diagnostic Lab Data:
CDC Split Type:

Write-up: Patient is not a patient of our clinic. When called to confirm appointment for second dose, we were notified patient was deceased.

VAERS ID: 1066344 (history)
Form: Version 2.0
Age: 88.0
Sex: Male
Location: Louisiana
Vaccinated: 2021-01-28
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9265 / 1	LA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: unknown
Current Illness: unknow
Preexisting Conditions: unknown
Allergies: unknown
Diagnostic Lab Data: N/A

CDC Split Type:

Write-up: This not a patient of our clinic. When called to confirm his second dose vaccine, is when we learned patient is deceased. Unable to reach anyone regarding date of death.

VAERS ID: 1068295 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Oklahoma
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-03

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

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Condition aggravated, Death, Illness, Malaise, SARS-CoV-2 test
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Lupus syndrome
Allergies:
Diagnostic Lab Data: Test Name: COVID-19; Test Result: Negative ; Comments: tested negative 2 times over the following 10 day while deteriorating.; Test Name: COVID-19; Test Result: Negative ; Comments: tested negative 2 times over the following 10 day while deteriorating.; Test Name: COVID-19; Test Result: Positive ; Comments: Was admitted and tested positive and put on ventilator
CDC Split Type: USPFIZER INC2021201566

Write-up: Was admitted and tested positive and put on ventilator; She felt slightly ill the day of vaccine; 2 days later patient become ill; tested negative 2 times over the following 10 day while deteriorating; Patient died 10 days later; This is a spontaneous report from a contactable consumer via Pfizer Sales Representative. This consumer (daughter) was reported for a female patient (mother). A 76-year-old female patient received first dose of bnt162b2 (Pfizer),

via an unspecified route of administration on an unspecified date at single dose for covid-19 immunization. Medical history included lupus from an unknown date and unknown if ongoing. The patient''s concomitant medications were not reported. The patient with lupus received 1st dose of vaccine. She felt slightly ill the day of vaccine. 2 days later patient become ill, tested negative 2 times over the following 10 day while deteriorating. Was admitted and tested positive and put on ventilator. Patient died 10 days later. Daughter thought she had COVID before vaccination. Event took place after use of product. The patient underwent lab tests and procedures which included COVID-19: negative (tested negative 2 times over the following 10 day while deteriorating), COVID-19: positive (Was admitted and tested positive and put on ventilator) all on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of other events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Patient died 10 days later

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VAERS ID: 1068307 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-03

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Haemorrhage
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021213418

Write-up: died; bled out; This is a spontaneous report from a contactable nurse. 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 medical history and concomitant medications were not reported. The patient had bled out on an unspecified date with outcome of unknown. The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Sender"s Comments: The limited information provided precludes a full clinical assessment of the case. As a cautionary measure and for reporting purposes, and assuming a drug-event temporal association, the Company cannot completely exclude a causal association between the reported events "bled out" and "died" (death of unknown cause) and BNT162B2 administration, until sufficient 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.,Linked Report(s) : US-PFIZER INC-2021210582 Same reporter/drug, different patient; Reported Cause(s) of Death: died

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VAERS ID: 1070005 (history)
Form: Version 2.0
Age: 85.0
Sex: Male
Location: Texas
Vaccinated: 2021-01-07
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: COVID-19, Death, SARS-CoV-2 test positive
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-03
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Hypertension, CAD, Hypothyroidism, Hypercholesterolemia
Preexisting Conditions: Hypertension, CAD, Hypothyroidism, Hypercholesterolemia
Allergies: NKA
Diagnostic Lab Data: Positive Covid Test

CDC Split Type:

Write-up: Patient contracted Covid some period of time after receiving the vaccine.

VAERS ID: 1070051 (history)
Form: Version 2.0
Age: 79.0
Sex: Female
Location: Texas
Vaccinated: 2021-01-06
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	025L20A / 1	AR / IM

Administered by: Private **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 2021-01-15
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Gabapentin, Pennsaid, Carbidopa-Levodopa
Current Illness: HTN, Aortic Valve Disorder, Coronary Atherosclerosis of Native Coronary Artery, Hyperlipidemia, Aortic aneurysm
Preexisting Conditions: HTN, Aortic Valve Disorder, Coronary Atherosclerosis of Native Coronary Artery, Hyperlipidemia, Aortic aneurysm
Allergies: Unknown
Diagnostic Lab Data: unknown
CDC Split Type:

Write-up: unknown

VAERS ID: 1070765 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021215418

Write-up: died 2 days after the second vaccine; This is a spontaneous report from a contactable consumer reporting for his/her father. An 87-year-old male patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number unknown) via unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient previously received the first dose of BNT162B2 on an unspecified date for COVID-19 immunisation and was fine. The patient died 2 days after the second vaccine. The reporter stated patient death due to the Pfizer Covid vaccine. The patient had autopsy. The outcome of event was fatal. Information about lot/batch number has been requested.; Reported Cause(s) of Death: died 2 days after the second vaccine

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VAERS ID: 1070769 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Diabetes; Heart disorder; Hospitalization; Pacemaker insertion (cardiac)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021224736

Write-up: then died within 24 hours afterwards; This is a spontaneous report from a contactable consumer. This consumer reported that a 74-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number unknown), via an unspecified route of administration an unspecified date at single dose for COVID-19 immunization. Concomitant medications were not reported. The patient with a host of health issues (heart issues/had a pace maker, diabetes, among others) was in a rehabilitation center following a hospital stay and was given the second dose of vaccine. It was reported that the patient died within 24 hours afterwards. It was not reported if an autopsy was performed. Information on batch/lot number was requested.; Reported Cause(s) of Death: then died within 24 hours afterwards

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VAERS ID: 1071300 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; A spontaneous report, was received from a consumer on a social media, concerning a 38-years-old female patient, unknown race and ethnicity, who was administered Moderna"s COVID-19 vaccine (mRNA-1273), and died. The patient"s medical history was not provided. No relevant concomitant medications were reported. On an unknown date, prior to the onset of the event, the patient received dose of mRNA-1273 (Lot number: unknown), for the prophylaxis of COVID-19 infection. On 17-Feb-2021, social media interaction was posted concerning a death of a patient on an unknown date after receiving Moderna vaccine. No treatment information was provided. Action taken with mRNA-1273 in response to the event was not applicable. The patient died on an unknown date. The cause of death was not provided. Plans for autopsy were not provided.; Reporter"s Comments: Very limited information regarding the event of death has been provided at this time. No further information will be available.; Sender"s Comments: MODERNA, INC.-MOD-2021-018380:Same reporter; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1073471 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Hospice care
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Was found deceased a little less than 12 hours following COVID vaccination; A spontaneous report was received from a reporter concerning a 96-year-old, male patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and was found deceased a little less than 12 hours following COVID vaccination. The patient"s medical history included hospice care. No relevant concomitant medications were reported. On an unknown date, prior to the onset of the event, the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. On an unknown date, the patient was found deceased a little less than 12 hours following COVID vaccination, and he had had some changes over the last two days. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not applicable. On an unknown date the patient died. The cause of death was unknown. Plans for an autopsy were not provided.; Reporter"s Comments: This case concerns a 96 year old male patient, who was on hospice care experienced a fatal event of death, after receiving mRNA- 1273 (Lot# Unknown). Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1076912 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-05

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?

Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021224962

Write-up: died from COVID after receiving the two doses of the vaccine; COVID; This is a spontaneous report from a contactable consumer report for a friend. A patient of unspecified age and gender received first dose and second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) both via an unspecified route of administration on unspecified dates at single doses for covid-19 immunisation. The patient"s medical history and concomitant medications were not reported. The consumer mentioned her friend died from COVID after receiving the two doses of the vaccine. The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: died from COVID after receiving the two doses of the vaccine

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VAERS ID: 1076917 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-05

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021229958

Write-up: one died after the vaccine; This is a spontaneous report from a Pfizer-sponsored program. This contactable consumer reported that a female patient of an unspecified age received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date at single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The reporter reported 3 females died post vaccination with the Pfizer-BioNTech COVID-19 vaccine. She explained one died after the vaccine on an unspecified date. She explained she had no additional details on the adverse event. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: one died after the vaccine

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VAERS ID: 1080335 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-08

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Anaphylaxis; A spontaneous report was received from a physician assistant concerning a patient of unspecified age and gender, who received Moderna"s COVID-19 vaccine (mRNA-1273) and experienced anaphylaxis. The patient"s medical history was not provided. No relevant Concomitant medications were reported. On an unknown date, prior to the onset of the event, the patient received their first of two planned doses of mRNA-1273 (Lot number: unknown) for prophylaxis of COVID-19 infection. On an unknown date, after receiving vaccine, the patient died due to anaphylaxis. No further details were available at the time of this report. Treatment for the event was not provided. Action taken with mRNA-1273 in response to the events was not applicable. The outcome of the event anaphylaxis was fatal .The patient died on an unspecified due to anaphylaxis. Autopsy details were not provided.; Reporter"s Comments: Very limited information regarding the event of anaphylaxis has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Anaphylaxis

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VAERS ID: 1083754 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Asymptomatic COVID-19, Drug ineffective, Systemic inflammatory response syndrome
SMQs:, Lack of efficacy/effect (narrow), Tumour lysis syndrome (broad), Sepsis (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021239063

Write-up: systemic inflammatory; asymptomatic COVID-19 infection; asymptomatic COVID-19 infection; This is a spontaneous report from a non-contactable consumer. A male patient of an unspecified age received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date at single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. On an unspecified date, the patient developed a devastating systemic inflammatory response within 36 hours of receiving his second Pfizer vaccine shot. Patient''s deadly inflammatory state followed his vaccination. Administration of the Pfizer vaccine ignited a deadly inflammatory response in patient''s body acutely, in a setting where he had a recent asymptomatic COVID-19 infection. The patient died on an unspecified date. It was not reported if an autopsy was performed. Outcome of all events were fatal. No follow-up attempts are possible, information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: asymptomatic COVID-19 infection; asymptomatic COVID-19 infection; systemic inflammatory

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VAERS ID: 1085865 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Ohio
Vaccinated: 2021-01-29
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-09

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	007M20A / 1	LA / IM

Administered by: Public **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-23
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Unknown
Current Illness:
Preexisting Conditions: unknown
Allergies: Unknown
Diagnostic Lab Data: Unknown
CDC Split Type:

Write-up: DEATH 2/23/21

VAERS ID: 1090182 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Tennessee
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-11

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Passed away; A spontaneous report was received from a consumer concerning a male patient of unknown age, who was received Moderna''s COVID-19 vaccine (mRNA-1273) and died. The patient''s medical history was not provided. No concomitant medications were reported. On an unknown date, prior to the onset of event, the patient received their first of two planned doses of mRNA-1273 (Lot number: unknown) intramuscularly for prophylaxis of COVID-19 infection. It was reported by the patient''s wife that the patient died. She was calling to cancel his second dose of mRNA-1273. No additional details, including the date of death, were reported. Treatment information was not provided. Action taken with the mRNA-1273 was not applicable as the patient died. The cause of death was not provided. Plans for an autopsy were not provided.; Reporter''s Comments: Very limited information regarding this event has been provided at this time. No further information is expected.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1094259 (history)
Form: Version 2.0
Age: 65.0
Sex: Male
Location: Unknown
Vaccinated: 2021-01-30
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL8982 / 1	RA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-26
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: unknown death

VAERS ID: 1100247 (history)
Form: Version 2.0
Age: 78.0
Sex: Female
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Discomfort, Loss of consciousness
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), 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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No recorded medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Unconsciousness; Feeling discomfort within a few minutes after the vaccination; A spontaneous report was received from a healthcare professional concerning a 78-year-old, female patient, who received Moderna"s COVID-19 Vaccine (mRNA-1273) and experienced of feeling discomfort within a few minutes after the vaccination, followed by unconsciousness and death. The patient"s medical history was not provided. No relevant concomitant medications were reported. On an unspecified date, prior to the onset of the event, patient received their dose of mRNA-1273 (Batch Number: Unknown) for prophylaxis of COVID-19 infection. On an unspecified date, the patient experienced feeling discomfort within a few minutes after the vaccination, followed by unconsciousness and death. Treatment information for event unconsciousness included resuscitate via cardiopulmonary resuscitation with no success. Action taken with mRNA-1273 in response to the events was not applicable. The

outcome of the event unconsciousness was fatal and patient died.; Reporter''s Comments: This is a case of death of a 78-year-old, female patient who experienced within a few minutes of product use who experienced feeling of discomfort and became unconscious and subsequently expired. Based on the current available information, a strong temporal association between the product use and onset of the events, a causal relationship with the events cannot be excluded. However, critical details such as the patient''s medical history and actual cause of death is lacking.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1106175 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Kentucky
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Sudden death
SMQs:, Torsade de pointes/QT prolongation (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: a friend died from the vaccine; A spontaneous report was received from a non health care professional, concerning her friend, who was administered Moderna''s COVID-19 vaccine (mRNA-1273) and died from the vaccine. The patient''s medical history,concomitant history, and lab data was not provided by the reporter. On an unknown date, the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. The patient experienced death after receiving the moderna mRNA1273 vaccine. No further details were provided. The action taken with the second dose of mRNA-

1273 in response to the event was not applicable. The outcome of the events, died from the vaccine, were considered as fatal.; Reporter"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1106349 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021259374

Write-up: Resulted in the death; This is a spontaneous report from a Pfizer sponsored program: A contactable consumer reporting on behalf of the sister reported that a female patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unknown date, at single dose, for COVID-19 immunisation. Medical history and concomitant medications were not reported. The patient experienced an adverse effect that resulted in death on an unspecified date. Cause of death was unknown. It was unknown if an autopsy was done. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Resulted in the death

VAERS ID: 1108473 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Arrhythmia, Cardiac failure congestive, Condition aggravated, Death, Diabetes mellitus, Fall
SMQs:, Cardiac failure (narrow), Hyperglycaemia/new onset diabetes mellitus (narrow), Accidents and injuries (narrow), Cardiomyopathy (broad), Cardiac arrhythmia terms, nonspecific (narrow), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-08
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: death Narrative: Patient received 1st dose of Moderna COVID-19 vaccine on 03/08/2021. Patient died 03/09/2021. Medical examiner received report that patient was alert before a fall on the night of 03/08/2021. Death certificate will report death likely due to arrhythmia due to underlying CHF. Contributing factors include diabetes.

VAERS ID: 1108474 (history)
Form: Version 2.0
Age:
Sex: Male

Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac failure, Cardiac failure congestive, Confusional state, Death, Dyspnoea, General physical health deterioration, Generalised oedema, Increased upper airway secretion, Insomnia, Nausea, Pulse absent, Respiratory arrest, Tachypnoea, Unresponsive to stimuli
SMQs:, Cardiac failure (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Angioedema (broad), Asthma/bronchospasm (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Dementia (broad), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (broad), Respiratory failure (narrow), Hypoglycaemia (broad), Infective pneumonia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-10
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: death Narrative: Patient was admitted to the ER on 12/26/20 with worsening shortness of breath and was admitted to acute care services. On 12/29/20, a hospice consult was placed for end stage CHF, EF 20%. On 12/30/20, he transferred to the facility and was a DNR. On 1/7/21, he was noted to have increased secretions in throat and was given atropine gtts sublingual and ondansetron for nausea. He also had issues with insomnia and was given trials of hydroxyzine, trazodone and melatonin. Lorazepam remained on profile as well as part of hospice care. On 1/9/21, he was noted to be more confused, tachypneic and had anasarca (furosemide was ordered). Later on that same day he began to decline rapidly to the point of

unresponsiveness other than to verbal stimuli and was determined to be imminent. On 1/10/21, he remained unresponsive and not able to tolerate oral meds. That same day at 1020 when nursing did rounds, he was found to be pulseless and without respirations. An autopsy was declined.

VAERS ID: 1107188 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Missouri
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-17

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: death after second dose; A spontaneous report was received from a consumer concerning a female patient of unknown age, who experienced severe symptoms post first dose of Moderna"s Covid-19 vaccine (mRNA-1273) and death shortly after her second dose. The patient"s medical history was not provided. Concomitant medication was not provided by the reporter. No lab data was not provided On unspecified date, the patient received her second of two planned doses of mRNA-1273 (Batch number not provided) intramuscularly for prophylaxis of COVID-19 infection. It was reported that shortly after receiving the second dose of mRNA-1273, the patient died. Treatment information was not provided/unknown. Action taken with the drug in response to the events is not applicable. The patient died on an unknown date. Plans for an autopsy were not reported; Reporter"s Comments: Very limited information regarding this event has been provided at this time. Further information has been

requested.; Reported Cause(s) of Death: death

VAERS ID: 1112825 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Arkansas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-18

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

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Abdominal pain, Arthralgia, Death, Epistaxis, Fatigue, Feeling abnormal, Mouth haemorrhage, Nausea, Pain, Pain in extremity, Pyrexia, Vomiting
SMQs:, Acute pancreatitis (broad), Haemorrhage terms (excl laboratory terms) (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Retroperitoneal fibrosis (broad), Dementia (broad), Oropharyngeal conditions (excl neoplasms, infections and allergies) (narrow), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Arthritis (broad), Tendinopathies and ligament disorders (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-17
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: None
CDC Split Type:

Write-up: Received vaccine on afternoon of 3/15/2021 and began experiencing nausea and vomiting, left arm and shoulder pain, fever, body aches, fatigue, and abdominal pain the morning of 3/16/20. He notified the pharmacy that administered the vaccine to him and they told him that some people have those symptoms with it. On 3/16/21 he went to bed around 11:30 PM feeling terrible and was found dead in his bed the next day (3/17/21) with dried blood coming out of his nose and mouth.

VAERS ID: 1114256 (history)

Form: Version 2.0
Age:
Sex: Male
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-19

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Aneurysm, Erythema
SMQs:, Anaphylactic reaction (broad), Hypersensitivity (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Aneurysm; red splotches on arm skin; A spontaneous report was received from a Consumer concerning a HCP, male patient of unknown age who received Moderna''s COVID-19 vaccine (mRNA-1273) and experienced developed red splotches on his arm skin and patient died from an aneurysm. The patient''s medical history was not provided. No concomitant medications were reported. On an unknown date, prior to the onset of the events, the patient received their unknown of the two planned doses of mRNA-1273 (lot/batch: unknown) via unknown route for prophylaxis of COVID-19 infection. On an unspecified date, after vaccination the patient had a reaction and developed red splotches on his arm skin. Then four to five days, later in the hospital the patient died from an aneurysm (Seriousness criteria: death). The cause of death was aneurysm. No autopsy details reported No Laboratory investigations were provided. No Treatment information was provided. Action taken with mRNA-1273 in response to the events was not applicable. The outcome of aneurysm was fatal and for erythema was unknown.; Reporter''s Comments: Very limited information regarding these events has been provided at this time. Further information has been requested; Reported Cause(s) of Death: Aneurysm

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VAERS ID: 1114383 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-19

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Liver disorder
SMQs:, Hepatic failure, fibrosis and cirrhosis and other liver damage-related conditions (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021284281

Write-up: The patient had liver problems and died; This is a spontaneous report from a contactable consumer. A 39-year-old female patient 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 medical history and concomitant medications were not reported. The patient had liver problems and died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: The patient had liver problems and died

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VAERS ID: 1114894 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Massachusetts

Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-19

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Product administered at inappropriate site, Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Systemic lupus erythematosus (broad), Drug abuse and dependence (broad), Medication errors (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Thrombocytopenia; Vaccinators are vaccinating very high up on the shoulder; A spontaneous report was received from a nurse concerning a patient, who received Moderna"s COVID-19 vaccine (mRNA-1273) and experienced Vaccinators are vaccinating very high up on the shoulder, thrombocytopenia. The patient"s medical history, was not provided by the reporter.No Concomitant medications were reported. On an unknown date, the patient received their planned dose of mRNA-1273 (Lot number: unknown) for prophylaxis of COVID-19 infection. The reporter stated that Vaccinators are vaccinating very high up on the shoulder. They need to go down on the deltoid. She also reported the event of thrombocytopenia of which one involved a doctor who died. Action taken with mRNA-1273 in response to the events was not applicable. The outcome of the event, vaccinators are vaccinating very high up on the shoulder was resolved . The outcome of the events thrombocytopenia is fatal.; Reporter"s Comments: This report refers to a patient who experienced non-serious of vaccinating very high up on the shoulder, (Vaccine administered at inappropriate site). There were no reported AEs associated with this case of vaccine administered at inappropriate site.; Reported Cause(s) of Death: Thrombocytopenia

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VAERS ID: 1116373 (history)
Form: Version 2.0
Age: 88.0

Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-19

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Asthenia, Death
SMQs:, Guillain-Barre syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No adverse event history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Passed away; so weak she was no longer able to use her walker and declined steadily each daythereafter; A spontaneous report was received from a consumer concerning a 88-year-old female patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and experienced being so weak she was no longer able to use her walker and declined steadily each day thereafter (asthenia) and passed away (death). The patient"s medical history was not provided. Concomitant product use was not provided by the reporter. The patient received their first of two planned doses of mRNA-1273 (Batch number not provided) on an unspecified date. On an unspecified date, approximately 15 days prior to the onset of the symptoms, the patient received their second of two planned doses of mRNA-1273 (Batch number not provided) intramuscularly for prophylaxis of COVID-19 infection. On an unspecified date, two days after their second dose of the vaccine, the patient was so weak that she was no longer able to use her walker and declined steadily each day thereafter. On an unspecified date, 17 days after their second dose of the vaccine, the patient passed away. Treatment information was not provided. The outcome of the event, asthenia, was unknown. The outcome of the event, passed away (death), was fatal. The cause of death was reported as unknown. Plans for an autopsy were not provided.; Reporter"s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1121585 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-22

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (no adverse event reported)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: died; A spontaneous report from was received from a Consumer concerning a female patient who received Moderna"s COVID-19 vaccine (mRNA-1273) and mentioned that a patient who had received the Moderna vaccine had died, and so they were concerned about getting the second dose. The patient"s medical history was not provided. No relevant concomitant medications were reported. On unknown date, prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. On an unknown date, The patient mentioned that a patient who had received the Moderna vaccine had died, and so they were concerned about getting the second dose. Laboratory details are not provided. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the events was considered Fatal.; Reporter"s Comments: This is a case of death of an unknown age female subject with unknown medical history, who died on an unknown day after receiving first dose of vaccine. Very limited information has been provided at this time. No follow up is possible.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1121622 (history)
Form: Version 2.0

Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-22

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021288934

Write-up: my friend just died after taking your shot; This is a spontaneous report from a Pfizer sponsored program. A non-contactable consumer reported a patient (friend) of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. His/her friend just died after taking your shot (And he/she know of more.) He/she will just keep his/her immune system high. They didn"t need a vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained; Reported Cause(s) of Death: my friend just died after taking your shot

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VAERS ID: 1121847 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: New York

Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-22

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Hepatic failure, Palpitations, Renal failure, Vomiting
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Hepatic failure, fibrosis and cirrhosis and other liver damage-related conditions (narrow), Acute pancreatitis (broad), Arrhythmia related investigations, signs and symptoms (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), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Chronic kidney disease (narrow), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (no relevant medical history reported)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Kidneys and liver shutting down; Kidneys and liver shutting down; Heart started racing 3 days later; Vomiting; A spontaneous report was received from a consumer concerning a 39-year-old, female patient, who received Moderna''s COVID-19 vaccine (mRNA-1273) and experienced heart racing, kidney failure, liver failure,vomiting and death. The patient''s medical history included Sjogren''s syndrome, Hashimoto''s, inflammatory joint disease, osteoporosis, osteoarthritis. She has history of fractured tail bone and leg in three places. Concomitant medications included D3 2 pills a day, folic acid, B12, and calcium citrate. She was allergic to 15 different antibiotics two of which she had anaphylaxis; had huge autoimmune issues; had had reaction to Pneumovax vaccine. On an unown date ,prior to the onset of the events, the patient received their second of two planned doses of mRNA-1273 intramuscularly for prophylaxis of COVID-19 infection. Approximately after three days of receiving second dose vaccine patient experienced heart racing and went to the emergency room .The patient started vomiting and developed a kidney and liver failure. And on the next day she died.Autopsy is doing and the results are not available at the time of report. Action taken with mRNA-1273 in response to the events was reported as not applicable. The events

heart racing, kidney failure, liver failure and death were considered serious and medically significant. The outcome of the events renal failure,hepatic failure,palpitations and vomiting was considered as Fatal.; Reporter''s Comments: This is a case of death in a 39year-old female subject with unknown medical history of anaphylactic reactions, allergic reactions to pneumovax vaccine, Sjogren''s syndrome, Hashimoto''s, inflammatory joint disease, osteoporosis, osteoarthritis, who died 4 days after receiving second dose of vaccine. Very limited information has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Renal failure; Hepatic failure

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VAERS ID: 1126281 (history)
Form: Version 2.0
Age: 93.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-03-16
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-23

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

Administered by: Public **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-18
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Asthma, AFib, Hyperlipidemia CHF, COPD
Allergies: Demerol, PCN
Diagnostic Lab Data: 0
CDC Split Type:

Write-up: unknown if related but resident Expired 3/18/21 48 hours after 2nd Covid vaccination Pfizer

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VAERS ID: 1126560 (history)

Form: Version 2.0
Age:
Sex: Male
Location: New Hampshire
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Chest pain, Death, 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? Yes
Date died: 0000-00-00
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 INC2021281269

Write-up: passed away; chest pain; trouble breathing; This is a spontaneous report from a contactable physician and from three non-contactable consumers from a Pfizer-sponsored program. A 66-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection; lot number and expiry date not reported), via an unspecified route of administration, on an unspecified date, as SINGLE DOSE for covid-19 immunisation. The patient had just taken the COVID-19 vaccine and he was hospitalized due to the effects. The patient was rushed to hospital with chest pains and was experiencing trouble breathing. The patient was in the ICU fighting the effects of the vaccine. The patient passed away four hours late on an unspecified date. The outcome of chest pains and trouble breathing was unknown. Cause of death was unknown. It was unknown if an autopsy was performed. Information on the lot/Batch number has been requested.; Sender''s Comments: Based on temporal association, the causal relationship between BNT162B2 and the events death, chest pain and dyspnea 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.; Reported Cause(s) of Death: passed away

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VAERS ID: 1130148 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Missouri
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Liver disorder
SMQs:, Hepatic failure, fibrosis and cirrhosis and other liver damage-related conditions (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: something wrong with the liver; A spontaneous report was received from a consumer concerning patients of unspecified age and gender who received Moderna''s COVID-19 vaccine (mRNA-1273) and were dying after second shot and something wrong with the liver (liver disorder). The patients'' medical histories were not provided. No relevant concomitant medications were reported. Dates of vaccination were not reported. The reported heard on the news that "people are dying after the second shot" and apparently one of the patients had something wrong with the liver after getting the shot. The reporter said the patients received Moderna''s mRNA-vaccine. On an unknown date, the patient experienced the event(s) people are dying after second shot, No treatment information was provided. The

action taken with the vaccine in response to the events was not applicable. The cause of death for "one of the patients was had something wrong with the liver".; Reporter"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested; Reported Cause(s) of Death: liver disorder; death unknown cause

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VAERS ID: 1133749 (history)
Form: Version 2.0
Age: 70.0
Sex: Male
Location: Unknown
Vaccinated: 2021-03-21
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EN6204 / 1	RA / IM

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-22
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: V-Fib, cardiac arrest Narrative: First COVID vaccine administered 2/25/21 with no noted reaction. Patient received his second COVID vaccine 3/21/21 at 1203. Notes in electronic medical record indicate in the morning of 3/22/21 he arrived at a hospital with ER this morning in V-Fib/cardiac arrest. Unclear of potential treatment that was administered at

outside facility. Time of Death was about 1330 on 3/22/21.

VAERS ID: 1148803 (history)
Form: Version 2.0
Age: 73.0
Sex: Male
Location: Unknown
Vaccinated: 2021-02-10
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL8982 / 2	- / -
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL0142 / 1	- / -

Administered by: Other **Purchased by:** ?
Symptoms: COVID-19, COVID-19 pneumonia, Death
SMQs:, Infective pneumonia (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-05
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-19/hospitalization/death Narrative: Patient received two doses of the Pfizer mRNA vaccine on 1/17/21 and 2/10/21. He was admitted on 3/5/21 with COVID-19 pneumonia which progressed to severe disease. Patient died on 3/21/21 attributed to COVID-19. Patient had underlying B cell follicular lymphoma, previously on chemotherapy. Reported does not feel patient''s hospitalization or death was attributable to the vaccines, however following FDA Emergency Use Authorization, "The vaccination provider is responsible for mandatory reporting of the following to the Vaccine Adverse Event Reporting System (VAERS): vaccine administration errors whether or not associated with an adverse event, serious adverse events* (irrespective of attribution to vaccination), cases of Multisystem Inflammatory Syndrome (MIS) in adults and children, and cases of COVID-19 that result in

hospitalization or death."

VAERS ID: 1133038 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest, Death, Malaise
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021296570

Write-up: died after receiving the first dose of a COVID-19 vaccine; cardiac arrest; fell ill; This is a spontaneous report from a non-contactable consumer reported for a patient (friend"s cousin). A male patient of an unspecified age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, batch/lot number and expiration date not provided), via an unspecified route of administration, on an unspecified date, at single dose, for COVID-19 immunization. Medical history included patient was infected with the virus in 2020 (reported as "over six months ago"), but not in the current time. Concomitant medications were not reported. The patient died after receiving the first dose of a COVID-19 vaccine on an unknown date. Patient felt ill that evening a few hours after receiving the shot, followed by cardiac

arrest. Patient was taken to the hospital, where he died the next day. The outcome of the event "unknown cause of death" was fatal, of other events was unknown. The cause of death was not reported. It was unknown if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: died after receiving the first dose of a COVID-19 vaccine

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VAERS ID: 1133040 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-25

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021297473

Write-up: died from heart attack; This is a spontaneous report from a contactable consumer via a Pfizer Sponsored reported for a male patient (Husband). A male patient of an unspecified age received first dose of COVID-19 vaccine (UNSPECIFIED TRADE NAME), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient died from heart attack exactly 1 week later after vaccine on an unspecified date. Never had any heart issues or anything. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: died

from heart attack

VAERS ID: 1135589 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021308003

Write-up: Death; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received bnt162b2 (BNT162B2), via an unspecified route of administration on an unspecified date (batch/lot number was not reported) as single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient got a steroid shot in between his 2 doses of the vaccine and he fell dead the next day. The reporter stated that the reporter didn''t know what manufacturer he received, but that he took steroids all the time. The patient died on an unspecified date. The cause of death was unknown. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1136163 (history)

Form: Version 2.0
Age: 88.0
Sex: Male
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EN6198 / 2	LA / SYR

Administered by: Public **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-24
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Keppra Potassium Triamterene Quetiapine
Current Illness:
Preexisting Conditions: Seizures Heart Dementia Stroke 12 yrs ago
Allergies: Penicillin
Diagnostic Lab Data:
CDC Split Type:

Write-up: Death

VAERS ID: 1140697 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-28

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

PFIZER/BIONTECH		

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021321350

Write-up: She passed one to two days after the shot; This is a spontaneous report from a contactable consumer. 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 single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. Patient died after receiving the COVID vaccine. She passed one to two days after the shot. It was not expected. Waiting on the autopsy. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: She passed one to two days after the shot

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VAERS ID: 1145721 (history)
Form: Version 1.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-30

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

Administered by: Unknown **Purchased by:** Unknown
Symptoms: COVID-19, Death
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: 2021028911

Write-up: COVID-19 INFECTION (COVID-19, COVID-19) This spontaneous report received from a consumer via a company representative and concerned a patient of unspecified age and sex. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. We are unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient developed covid-10 infection. On an unspecified date, the patient died from covid-19. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).

VAERS ID: 1146779 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Mississippi
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021321304

Write-up: passed away; This is a spontaneous report received from a contactable consumer reporting for a male patient of unspecified age that received second dose of BNT162B2 via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. It was reported that the patient passed away late Saturday night after receiving his second COVID-19 Vaccine on Friday. An autopsy was not performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained; Reported Cause(s) of Death: passed away

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VAERS ID: 1146785 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021321442

Write-up: She died in the bathroom on the second day.; This is a spontaneous report from a contactable consumer. This reporter reported similar events for 3 patients. This is 2nd of 3 patients. A female patient of an unspecified age received bnt162b2 (BNT162B2, PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient medical history was not reported. The patient"s concomitant medications were not reported. It is reported the patient died in upstate. Waiting on the autopsy. She died in the bathroom on the second day. It was not expected. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the event was fatal. Information on Lot/Batch has been requested.; Reported Cause(s) of Death: She died in the bathroom on the second day.

VAERS ID: 1149905 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; A spontaneous report was received from a Consumer concerning a patient where age and gender unspecified who received Moderna''s COVID-19 vaccine (mRNA-1273) and experienced event Death. The patient''s medical history was not provided. No relevant concomitant medications were reported. On unknown date, prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (lot/batch: unknown) in the Anatomical location unspecified for prophylaxis of COVID-19 infection. On unknown date, The patient experienced the event Death. Laboratory details were not provided. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the event was Fatal. The reporter assessed the event Death related to the study drug was unknown.; Reporter''s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1153088 (history)
Form: Version 2.0
Age:
Sex: Female
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-03-31

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210331140

Write-up: MAJOR HEART ATTACK; This spontaneous report received from a consumer concerned a 50 year old female. The patient''s weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, start therapy date were not reported for

prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, she had major heart attack and died. It was unknown if an autopsy was performed. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The reporter and company provided causality between Covid-19 vaccine ad26.cov2.s and major heart attack as possible. This report was serious (Death).; Sender''s Comments: - covid-19 vaccine ad26.cov2.s -Heart attack. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: MAJOR HEART ATTACK

VAERS ID: 1153902 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Mississippi
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Pain in extremity
SMQs:, Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210353764

Write-up: DIED; SORE ARM/ARM PAIN; This spontaneous report received from a consumer concerned a female of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient

had a sore arm/arm pain, and later on died. On an unspecified date, the patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died on an unspecified date, and the outcome of sore arm/arm pain was not reported. This report was serious (Death). This case, from the same reporter is linked to 20210329044.; Sender''s Comments: V0: 20210353764 - COVID-19 VACCINE AD26.COV2.S - Died. This event is considered Unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1153994 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died from the Moderna vaccine; A spontaneous report was received from a consumer concerning a patient who received Moderna''s COVID-19 vaccine (mRNA-1273) and died from the Moderna vaccine/death. The patient''s medical history was not provided. No concomitant product use was reported. On unknown date, the patient received their first of two planned doses of mRNA-1273 (Lot number: not provided) intramuscularly for prophylaxis of COVID-19 infection. On unknown date, the patient died after receiving the Moderna vaccine

(Seriousness criteria: death, medically significant). The date of death was unknown. Autopsy results not provided. Treatment information was not provided. Action taken with mRNA-1273 in response to the events was not applicable. The outcome of the event, died from the Moderna vaccine/death, was considered fatal. .; Reporter"s Comments: Very limited information regarding the event of death has been provided at this time. No further information has been requested.; Reported Cause(s) of Death: Died from the Moderna vaccine

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VAERS ID: 1154126 (history)
Form: Version 2.0
Age: 33.0
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Diabetic ketoacidosis
SMQs:, Hyperglycaemia/new onset diabetes mellitus (narrow), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-21
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Gastroparesis (Verbatim: Gastroparesis); Type I diabetes mellitus (Verbatim: Type 1 Diabetes); Ulcer (Verbatim: Ulcer)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021331199

Write-up: Diabetic ketoacidosis; This is a spontaneous report from a contactable consumer. A contactable consumer reported for her nephew. A 33-years-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (at the age of 33-years-old) at single dose for covid-19 immunisation. Medical history included type 1 diabetes mellitus from an unknown date and unknown if ongoing, impaired gastric emptying from an unknown date and unknown if

ongoing, ulcer from an unknown date and unknown if ongoing. The patient"s concomitant medications were not reported. The patient experienced diabetic ketoacidosis on an unspecified date. The patient died on 21Mar2021 due to diabetic ketoacidosis. An autopsy was not performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Diabetic ketoacidosis

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VAERS ID: 1155738 (history)
Form: Version 2.0
Age: 54.0
Sex: Male
Location: Oregon
Vaccinated: 2021-02-19
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	693894 / 2	LA / SYR

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-19
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Gabapentin
Current Illness: None
Preexisting Conditions: Lumbar fusion
Allergies: None
Diagnostic Lab Data:
CDC Split Type:

Write-up: Death

VAERS ID: 1157484 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-01

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

Administered by: Other **Purchased by:** ?
Symptoms: COVID-19
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210328911

Write-up: COVID-19 INFECTION; This spontaneous report received from a consumer via a company representative and concerned a patient of unspecified age and sex. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. We are unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient developed covid-19 infection. On an unspecified date, the patient died from covid-19. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: A patient of unknown age and gender died an unknown time after receiving Janssen COVID-19 Vaccine Ad26.COV2.S (suspension for injection, route of administration not reported) for prophylactic vaccination. Medical history, concomitant medications, and details of the event were not reported. It was unknown if an autopsy was performed. This case has insufficient information to make a meaningful medical assessment.; Reported Cause(s) of Death: COVID-19

VAERS ID: 1157515 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-04-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Unknown cause of death; Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210350272

Write-up: DEATH; This spontaneous report received from a consumer concerned multiple patients. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported, per procedure no follow up will be requested for this case. No concomitant medications were reported. The reported called for an agency, during call she had mentioned "several deaths were reported with the vaccine". She read it on a website. The cause of the deaths was not reported. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: 20210350272-covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

VAERS ID: 1161961 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-04-02

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210356723

Write-up: DEATH; This spontaneous report received from a patient via a company representative concerned a patient of unspecified age and sex. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin)total 1 dose, start therapy date were not reported for prophylactic vaccination. The Batch number was not reported. The Company is unable to perform follow-up to request batch/lot number. No concomitant medications were reported. On an unspecified date, the patient experienced death. On an unspecified date, the patient died from unknown cause of death. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: 20210356723- Covid-19 vaccine ad26.cov2.s-Death. This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1162137 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-02

Vaccination / Manufacturer	Lot /	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021340045

Write-up: when the patient got really bad the patient died; This is a spontaneous report from a contactable consumer. A 92-year-old patient of an unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. The patient experienced when the patient got really bad the patient died. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: when the patient got really bad the patient died

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VAERS ID: 1166023 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-04

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

Administered by: Unknown **Purchased by:** ?

Symptoms: COVID-19, Completed suicide, Drug ineffective, Tinnitus
SMQs:, Lack of efficacy/effect (narrow), Suicide/self-injury (narrow), Hearing impairment (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021344589

Write-up: killed himself; tinnitus; contracted the virus; contracted the virus; This is a spontaneous report from a non-contactable nurse. A male patient of an unspecified age received BNT162B2 (Pfizer-BioNTech COVID-19 vaccine, lot number and expiry date unknown), via an unspecified route of administration on an unspecified date as a single dose for COVID-19 immunization. Medical history included COVID-19 on an unknown date. The patient''s concomitant medications were not reported. It was reported that the patient, contracted the virus and developed tinnitus after receiving the vaccine and killed himself. The outcome of the event tinnitus and contracted the virus was unknown. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow up attempts are possible; information about lot number/batch number cannot be obtained.; Sender''s Comments: Based on the information currently available, a causal association between the reported event tinnitus and BNT162B2 cannot be excluded. Drug ineffective depends on many fators including pharmacokinetics, patient general health condition and immunity system function. However on conservative basis, the possible causality cannot be excluded. The event "killed himself" is not related to 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 RAs, Ethics Committees, and investigators, as appropriate. ; Reported Cause(s) of Death: killed himself

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VAERS ID: 1166026 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-04-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Diarrhoea haemorrhagic, Haemoglobin, Platelet count, Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Pseudomembranous colitis (broad), Gastrointestinal haemorrhage (narrow), Ischaemic colitis (broad), Noninfectious diarrhoea (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Diverticulosis; Hypothyroidism; Transcatheter aortic valve implantation (recent)
Allergies:
Diagnostic Lab Data: Test Name: hemoglobin; Result Unstructured Data: Test Result:8.7 g/dl; Test Name: platelets; Result Unstructured Data: Test Result:60 x10 9/l; Test Name: platelets; Result Unstructured Data: Test Result:improvement to 101 x10 9/l; Comments: was discharged 5 days later
CDC Split Type: USPFIZER INC2021353725

Write-up: hemoglobin 8.7 g/dL and platelets 60*109/L; bloody diarrhea; This is a literature-spontaneous report based on the journal following Pfizer and Moderna SARS-CoV-2 vaccination. The same author reported different events for 4 patients received Pfizer vaccine. This is the 1st of 4 reports. The reports describing 19 of 20 patients with thrombocytopenia following vaccination Pfizer and Moderna SARS-CoV-2 vaccination, included age (range 22-73 years old; median 41 years) and gender (11 females and 8 males). Nine received the Pfizer vaccine and 11 received the Moderna vaccine. All 20 patients were hospitalized and most patients presented with petechiae, bruising or mucosal bleeding (gingival, vaginal, epistaxis) with onset of symptoms between 1-23 days (median 5 days) post vaccination. Platelet counts at presentation were available for all 20 cases with the majority being at or below 10*109/L (range 1-36*109/L; median 2*109/L). This case refers to an 80-year-old male patient with multiple medical problems including recent transcatheter aortic valve replacement, hypothyroidism, and diverticulosis presented 6 days after the Pfizer vaccine with bloody diarrhea, hemoglobin 8.7 g/dL and platelets 60*109/L. He received several units of packed red blood cells and two units of platelets with improvement to 101*109/L and was discharged 5 days later. There were a handful of reports with minimal additional details alluding to a male who passed away in Dec2020 from brain hemorrhage following the Pfizer

vaccine-these could be describing the index patient. There were a handful of reports with minimal additional details alluding to a male who passed away in December from brain hemorrhage following the Pfizer vaccine-these could be describing the index patient. In summary, we cannot exclude the possibility that the Pfizer and Moderna vaccines have the potential to trigger de novo ITP (including clinically undiagnosed cases), albeit very rarely. Distinguishing vaccine-induced ITP from coincidental ITP presenting soon after vaccination is impossible at this time. No follow up attempts are needed; information about lot/batch number cannot be obtained. No further information is expected.; Sender"s Comments: Based on the information currently provided (plausible time association), it cannot be fully excluded that the SARS-CoV-2 vaccination might have played a contributory role in triggering the occurrence of thrombocytopenia complicated by bloody diarrhea leading to blood loss requiring blood products infusion. 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) : US-PFIZER INC-2021353705 same reporter, different patient, same suspect , different event;US-PFIZER INC-2021014534 same reporter, different patient, same suspect , different event;US-PFIZER INC-2021344239 same reporter, different patient, same suspect , different event; Reported Cause(s) of Death: passed away in December from brain hemorrhage

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VAERS ID: 1168960 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-05

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown

Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210350465

Write-up: GOT THE VACCINE AND DIED 24 HOURS LATER; This spontaneous report received from a consumer concerned a patient of unspecified age and sex. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient died 24 hours later vaccination. Cause of death was not reported. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0:20210350465-covid-19 vaccine ad26.cov2.s-Got the vaccine and died 24 hours later. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s); Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1168970 (history)
Form: Version 2.0
Age: 95.0
Sex: Female
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-05

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

Administered by: Other **Purchased by:** ?
Symptoms: Atrial fibrillation, Death, Dysphagia, Hypotension, Peripheral embolism
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Supraventricular tachyarrhythmias (narrow), Embolic and thrombotic events, arterial (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), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210400509

Write-up: ATRIAL FIBRILLATION; INABILITY TO SWALLOW; BLOOD CLOT IN RIGHT ARM; LOW BLOOD PRESSURE; DEATH 4 DAYS AFTER RECEIVING VACCINE; This spontaneous report received from a vaccine facility via a company representative concerned a 95-year-old female. The patient''s height, and weight were not reported. The patient''s concurrent conditions included atrial fibrillation. The patient received COVID-19 VACCINE AD26.COV2.S (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered on 2021 for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported and has been requested. It was reported that on an unspecified date in 2021 the patient received Janssen Covid-19 Vaccine and within 6hrs she had a major atrial fibrillation episode, then several the following day. The next day, she lost her ability to swallow. Two days later she was on oxygen. Three days later she developed a blood clot in her right arm, was still on oxygen and blood pressure was falling. On an unspecified date, the patient died 4 days after receiving vaccine. The action taken with COVID-19 VACCINE AD26.COV2.S was not applicable. The patient died 4 days after receiving vaccine on an unspecified date, and the outcome of atrial fibrillation, inability to swallow, blood clot in right arm and low blood pressure was not reported. This report was serious (Death, Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender''s Comments: V0: 20210400509: This spontaneous report received from a vaccine facility via a company representative involved a 95-year-old female with the past medical history remarkable for atrial fibrillation who received the Janssen COVID-19 Vaccine for prevention of COVID-19 infection and within 6hrs had a major atrial fibrillation episode. No concomitant medications were reported. The next day, she lost her ability to swallow. Two days later she was on oxygen. Three days later she developed a blood clot in her right arm, was still on oxygen and blood pressure was falling. On an unspecified date, the patient died 4 days after receiving vaccine. No information was provided regarding the cause of death. Considering the patient''s past medical history of atrial fibrillation, the causality for the event of atrial fibrillation, as well the consequent events is assessed not related to the Janssen COVID-19 Vaccine.; Reported Cause(s) of Death: DEATH 4 DAYS AFTER RECEIVING VACCINE

VAERS ID: 1169011 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-05

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

Administered by: Unknown **Purchased by:** ?

Symptoms: COVID-19
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: one friend got COVID after the shot and has passed away; A spontaneous report was received from a consumer concerning a female patient of an unknown age who received Moderna"s COVID-19 vaccine and got COVID after the shot and has died/COVID-19. The patient"s medical history was not provided. Concomitant product use was not provided by reporter. On an unknown date the patient received their first of two planned doses of mRNA-1273 (Batch number not provided) for prophylaxis of COVID-19 infection. Reporter mentioned that her friend got COVID after the shot and has died. No treatment information was provided. Action taken with the second dose of mRNA-1273 in response to the event death is not applicable. The patient died on an unknown date. The cause of death was not provided. Plans for an autopsy were not provided.; Reporter"s Comments: A case of death of female patient of an unknown age who developed COVI on an unknown date post mRNA-1273 (vaccination (lot unknown) and died. Very limited information regarding this event/s has been provided at this time. However, based on the known etiology of COVID and the established profile of mRNA-1273, the event is assessed as unlikely related to mRNA-1273 administration. Further information has been requested.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1173591 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-06

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

Administered by: Unknown **Purchased by:** ?

Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No medical history reported)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; A spontaneus report was received from a consumer concerning a male patient of an unknown age who received Moderna''s COVID-19 vaccine (mRNA-1273) and died. The patient''s medical history was not provided. No relevant concomitant medications were reported. On an unknown date, prior to onset of the event, the patient received their dose of mRNA-1273 (lot/batch number: unknown) for prophylaxis of COVID-19 infection. On an unknown date, the patient died. Action taken with mRNA-1273 in response to the event was not applicable. The patient died on an unspecified date. The cause of death was unknown. Plans for an autopsy were unknown.; Reporter''s Comments: Very limited information regarding the event has been provided at this time and is insufficient for causality assessment. The cause of death was unknown. Plans for an autopsy were unknown. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1174358 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021352130

Write-up: Pfizer vaccination killed patient; This is a spontaneous report from a Pfizer-sponsored program, Corporate (Pfizer) Social Media Platforms. A contactable consumer reported for mother (patient) that a female patient of unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot/batch number and expiration date not provided), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The reporter reported that Pfizer vaccination killed reporter''s mom (patient) on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Pfizer vaccination killed patient

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VAERS ID: 1178144 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Oregon
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-07

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

Administered by: Other **Purchased by:** ?
Symptoms: Drug interaction
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: RISPERDAL
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210403502

Write-up: DIED FROM THE INTERACTION OF THE TWO PRODUCTS RISPERDAL AND COVID VACCINE; This spontaneous report received from a consumer who had heard that two people who took Risperdal and got a COVID vaccine died from the interaction of the two products. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. The patient received risperidone (form of admin, route of admin, and batch number were not reported) dose, frequency, and therapy dates were not reported for an unspecified indication. No concomitant medications were reported. On an unspecified date, consumer heard that a few people who took Risperdal and got a COVID vaccine died from the interaction of the two products. The action taken with covid-19 vaccine ad26.cov2.s, and risperidone was not applicable. This report was serious (Death).; Sender"s Comments: A report received from a consumer who had heard that "a few people who took Risperdal and got a COVID vaccine died from the interaction of the two products." The patients past medical history, concomitant medications were not reported. COVID-19 vaccine ad26.cov2.s date and dose administered were not reported. Risperidone dose, frequency, therapy dates and indication were not reported. There is insufficient information provided in this case to make a meaningful medical assessment.

VAERS ID: 1178152 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-07

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

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

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210403505

Write-up: HEART ATTACK; This spontaneous report received from a consumer via a company concerned a 40 year old male. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: UNKNOWN) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow up to request batch/lot number. No concomitant medications were reported. About 7 days after receiving the vaccine, the patient went to hospital regarding shortness of breath. On an unspecified date, the patient experienced heart attack and was hospitalized (date unspecified) and was later sent to intensive care unit (ICU). On an unspecified date, the patient died from heart attack. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Hospitalization Caused / Prolonged).; Sender''s Comments: 20210403505 -Covid-19 vaccine ad26.cov2.s -Heart attack. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: HEART ATTACK

VAERS ID: 1178296 (history)
Form: Version 2.0
Age:
Sex: Male
Location: North Carolina
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest

SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021347629

Write-up: cardiac arrest; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient experienced cardiac arrest 20 hours after receiving BNT162B2 on an unspecified date. The patient passed away due to cardiac arrest. The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: cardiac arrest

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VAERS ID: 1178308 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Washington
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021371687

Write-up: two deaths in her state from vaccine; This is a spontaneous report received from a contactable consumer. This consumer reported similar events for two patients. This the first of two reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number not provided), on unspecified date at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. Reporter was concerned because there had been two deaths in her state from vaccine and thought Pfizer might be curious in how their product was being handled. The patient died on unspecified date. It was unknown if autopsy was performed. The outcome of the event was fatal. Information on lot/batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021371688 same reporter/event, different patient; Reported Cause(s) of Death: two deaths in her state from vaccine

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VAERS ID: 1182765 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cardiovascular disease, unspecified
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021352646

Write-up: died approximately one week after receiving the Pfizer vaccine; This is a spontaneous report from a non-contactable consumer. A 94-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not reported), via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunization. Medical history included cardiovascular disease and other conditions (could not name). The patient"s concomitant medications were not reported. The patient was ill but was able to walk around on own and found pretty weak for age. It was reported that the patient died approximately one week after receiving the vaccine on an unspecified date. It was unknown if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: died approximately one week after receiving the Pfizer vaccine

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VAERS ID: 1192044 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-10

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021369331

Write-up: died; This is a spontaneous report from a contactable consumer. A 30-year-old female patient received her first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiry date were not reported), via an unspecified route of administration on an unspecified date as a single dose for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. It was reported that the patient who was completely healthy died on an unspecified date. Reporter stated that she had "great concerns about the Pfizer COVID 19 vaccine and actually all three of the vaccines on the market. She had been so distraught over the last couple of weeks since she found out that they are wanting to start experimentations on babies and that the babies can not handle this. She wanted to ask someone how does this fit in with the Nuremberg code since the babies have no awareness of what is going on since they are 6 months or less? She stated experiments should be based on animal experimentation but by the way all the companies skip animal testing. 15 years ago all the animals tested for the coronavirus vaccine died. The 30 year old mother that died bothered her completely and she never even met that person. The reporter stated people should have survived and that people do not need vaccine junk science and that it is making them into some operating system. She stated the vaccine is not helping and this was created just to create more fear." It was not reported if an autopsy was performed. Information about lot/batch number has been requested. ; Reported Cause(s) of Death: Died

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VAERS ID: 1192062 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Washington
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-10

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021371688

Write-up: death; This is a spontaneous report received from a contactable consumer. This consumer reported similar events for two patients. This the second of two reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number not provided) via an unspecified route of administration, on unspecified date at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient experienced death on an unspecified date. It was unknown if autopsy was performed. Information on lot/batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021371687 same reporter/event, different patient; Reported Cause(s) of Death: death

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VAERS ID: 1192078 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Washington
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cancer
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021372641

Write-up: Passed away; This is a spontaneous report from a contactable consumer received via a Pfizer-sponsored program, Pfizer RXPathways. A male patient of an unspecified age received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunisation. Medical history included stage 4 cancer. The patient"s concomitant medications were not reported. The patient passed away on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Passed away

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VAERS ID: 1192117 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-10

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebral haemorrhage
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhagic central nervous system vascular conditions (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021385762

Write-up: Brain bleed; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender (physician) received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date (at an unspecified age) at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The consumer reported that a perfectly healthy physician who died after receiving the vaccine actually dying from a brain bleed. The outcome of the event was fatal. The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Brain bleed

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VAERS ID: 1194127 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-11

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:

CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Died) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient''s past medical history included No adverse event (No reported medical history). On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product use was not provided by the reporter. No treatment information was provided. Very limited information regarding the event of death has been provided at this time. Further information has been requested.; Sender''s Comments: Very limited information regarding the event of death has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1205264 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Connecticut
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-13

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Malaise
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021390868

Write-up: person didn''t get the second shot, got sick and died; person didn''t get the second shot, got sick and died; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), first dose via an unspecified route of administration on an unspecified date (batch/lot number and expiration date was not reported) as single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. The consumer stated that they had a case where a person didn''t get the second shot, got sick and died (pending confirmation). The patient died on an unspecified date. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: person didn''t got the second shot got sick and died

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VAERS ID: 1207372 (history)
Form: Version 2.0
Age: 90.0
Sex: Female
Location: Puerto Rico
Vaccinated: 2021-01-13
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-14

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	032H20A / 1	UN / IM

Administered by: Senior Living **Purchased by:** ?
Symptoms: Death, Hypotension
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-01-21
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Probiotics and Omeprazole
Current Illness: None reported
Preexisting Conditions: Respiratory problems, in hospice
Allergies: None reported
Diagnostic Lab Data: None reported
CDC Split Type:

Write-up: On 1/21/2021 Caregivers proceed to go on rounds taking vitals and found she was with low blood pressure. They call Hospice and they certify that she had passed away.

VAERS ID: 1216626 (history)

Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-16

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	RA / IM

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: None stated.

VAERS ID: 1218468 (history)
Form: Version 2.0
Age: 82.0
Sex: Male
Location: Kentucky
Vaccinated: 2021-03-18
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-16

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	044A21A / 1	RA / IM

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

Symptoms: Death, Myocardial infarction
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-04-05
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"s daughter called and reported he was hospitalized for a heart attack and passed away on 4/5/2021.

VAERS ID: 1219067 (history)
Form: Version 2.0
Age: 64.0
Sex: Male
Location: Florida
Vaccinated: 2021-03-20
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-16

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	1805029 / 1	AR / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Death, Dyslipidaemia, Infarction, Surgery, Syncope
SMQs:, Torsade de pointes/QT prolongation (broad), Dyslipidaemia (narrow), Myocardial infarction (broad), Arrhythmia related investigations, signs and symptoms (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Cardiomyopathy (broad), Lipodystrophy (broad), Hypotonic-hyporesponsive episode (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-24
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Patient had been off of his statin medication of 2 weeks according to the daughter
Current Illness:
Preexisting Conditions: hyperlipidemia
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: 03/24/21, patient hotel , patient fainted , tried to revive by family , ems was called , patient underwent surgery , patient did not survive surgery , death certificate was for acute infarction and dyslipidemia , patient passed

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VAERS ID: 1220913 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 2021-02-03
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-16

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	012M20A / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Pancreatic carcinoma, Vaccination site pain
SMQs:, Non-haematological malignant tumours (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-23
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cancer (NOS) (for 30 years); Diabetes; Pancreatic cancer
Allergies:
Diagnostic Lab Data:

CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Husband died because of terminal pancreatic cancer; Sore arm; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PANCREATIC CARCINOMA (Husband died because of terminal pancreatic cancer) in a 74-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 012M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient"s past medical history included Diabetes. Concurrent medical conditions included Pancreatic cancer and Cancer (NOS) (for 30 years). On 03-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced PANCREATIC CARCINOMA (Husband died because of terminal pancreatic cancer) (seriousness criteria death and medically significant) and VACCINATION SITE PAIN (Sore arm). On 23-Feb-2021, VACCINATION SITE PAIN (Sore arm) outcome was unknown. The patient died on 23-Feb-2021. The reported cause of death was Pancreatic cancer. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. He was on loads of different medication for cancer which the wife did not know about. Treatment information was not provided. As per the patient"s wife on 23Feb2021, her husband died because of terminal pancreatic cancer which she thinks is not related to the vaccine This case was linked to US-MODERNATX, INC.-MOD-2021-069647 (Linked Report).; Sender"s Comments: Based on the current available information and the temporal association between the product use and the start date of the event a causal relationship cannot be ruled out. US-MODERNATX, INC.-MOD-2021-069647:; Reported Cause(s) of Death: Pancreatic cancer

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VAERS ID: 1220979 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Death; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient"s past medical history included No adverse event. Concurrent medical conditions included No adverse event. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant medication list was not provided. Treatment information was not provided.; Sender"s Comments: Limited information regarding the event has been provided at this time and a causal relationship cannot be excluded; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1221166 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Indiana
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-16

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Transplant
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: No adverse event (No reported medical history)

Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: needed kidney and/or bladder transplant; within 30 hs after second shot, she died; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of DEATH (within 30 hs after second shot, she died) and TRANSPLANT (needed kidney and/or bladder transplant) in a 39-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient"s past medical history included No adverse event (No reported medical history). On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced DEATH (within 30 hs after second shot, she died) (seriousness criterion death) and TRANSPLANT (needed kidney and/or bladder transplant) (seriousness criteria death, hospitalization and medically significant). The cause of death was not reported. It is unknown if an autopsy was performed. The reporter does not remember if this was from Pfizer or Moderna. Concomitant product us e was not provided. Treatment information was unknown. This case concerns death of a 30 year old female 30 minutes after the administration of the mRNA-1273 vaccine. Patient is reported to have no underlying issue. Based on the current available information which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, causal relationship cannot be excluded. Reporter did not allow further contact; Reporter"s Comments: Th; Sender"s Comments: This case concerns death of a 30 year old female 30 minutes after the administration of the mRNA-1273 vaccine. Patient is reported to have no underlying issue. Based on the current available information which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, causal relationship cannot be excluded.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1224899 (history)
Form: Version 2.0
Age: 94.0
Sex: Female
Location: Illinois
Vaccinated: 2021-03-23
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021389796

Write-up: passed away; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that a 94-year-old female patient (reporter''s mother) received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 23Mar2021 (Batch/Lot number was not reported) as a single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient passed away on an unspecified date. It was unknown if an autopsy was performed. The reporter said that patient died because she is 94 years old and not with the vaccine. Information on the lot/batch number has been requested.; Sender''s Comments: Based on the available information, this female patient death is attributed to her advanced age (94 years old) and assessed unrelated to BNT162B2 vaccine.; Reported Cause(s) of Death: passed away

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VAERS ID: 1227282 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Georgia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021404175

Write-up: I know a physician in died afterwards; This is a spontaneous report from a Pfizer-sponsored programs 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 (Lot number was not reported) as single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient knows a physician and died afterwards on an unspecified date. It was unknown if an autopsy was performed. Information about lot/batch number has been requested.; Reported Cause(s) of Death: I know a physician in (name) died afterwards

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VAERS ID: 1227286 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021416933

Write-up: within two weeks of receiving the vaccine the patient died; This is a spontaneous report based on information received by Pfizer from Merck & Co., Inc. A contactable Consumer reported for a male patient of an unspecified age that received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient medical history and the patient''s concomitant medications were not reported. The patient died within two weeks of receiving the vaccine on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible, information about lot/batch number cannot been obtained.; Reported Cause(s) of Death: within two weeks of receiving the vaccine both family members died

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VAERS ID: 1227287 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021416943

Write-up: within two weeks of receiving the vaccine both family members died; This is a spontaneous report based on information received by Pfizer (Case Number: 01903495). A contactable Consumer reported for two family members. A male patient of an unspecified age received bnt162b2 (BNT162B2), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient died on an unspecified date within two weeks of receiving the vaccine . It was not reported if an autopsy was performed. No follow-up attempts are possible, information about lot/batch number cannot been obtained.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1227818 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Clot blood (She was prone to blood clots her entire life, but managed it.)
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210420170

Write-up: BLOOD CLOT; This spontaneous report received from a consumer via media by a company representative concerned a female of unspecified age. The patient"s height, and weight were not reported. The patient was prone to blood clots her entire life, but managed it. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration and batch number were not reported) dose (1 total), start therapy date were not reported, for prophylactic vaccination. The batch number was not reported. Per procedure, no

follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient developed a blood clot. A day after the vaccination, the patient died from the blood clot. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: This anecdotal report from media involves a female patient of unspecified age who was prone to blood clots her entire life and on an unspecified date developed a blood clot and died from the blood clot a day after received the Janssen COVID-19 Vaccine Ad26.COV2. Concomitant medications, and details of the event were not reported. This case has insufficient information to make a meaningful medical assessment. The case will be assessed further when additional information is received.; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1227922 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Exposure during pregnancy, Skeletal injury, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad), Accidents and injuries (narrow), Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Gravida/Para: 1/1. The patient was 4 weeks post-partum.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210427241

Write-up: BLOOD CLOT; BROKE TAIL BONE; VACCINE EXPOSURE DURING PREGNANCY; This spontaneous pregnancy report was received from a pharmacist via a

company representative, and concerned an approximately 40 year old female. The patient''s height, weight, and medical history were not reported. The patient received Covid-19 vaccine Ad26.COV2.S (suspension for injection, route of administration not reported, batch number: unknown) dose and vaccination site were not reported, administered in 2021 for prophylactic vaccination. No concomitant medications were reported. In 2021, the patient experienced vaccine exposure during pregnancy. The date of the patient''s last menstrual period and expected delivery date were not provided. In 2021, the patient experienced broke tail bone during labor and gave birth (live birth). On an unspecified date in 2021, the patient experienced a blood clot and died. It was noted that she was at high risk for clots because she was 4 weeks post partum (gravida 1, para 1). Action taken with Covid-19 vaccine Ad26.COV2.S was not applicable. The patient died of a blood clot and broke tail bone in 2021; the outcome of vaccine exposure during pregnancy was not reported. It was unspecified if an autopsy was performed. This report was serious (Death). This case, from the same reporter is linked to 20210430297.; Sender''s Comments: V0: The case concerns a pregnant female subject around age of 40, who developed thrombosis, skeletal injury and exposure during pregnancy an unspecified time after Janssen COVID-19 vaccine was administered intramuscularly for prevention of symptomatic SARS-CoV-2 virus infection. The subject''s past medial history, last menstrual period, estimated date of delivery and concomitant medications were not provided. Per the reporter (pharmacist) the patient was at a high risk for blood clots because she was 4 weeks post-partum. The patient broke her tail bone during the labor, gave a birth, and later died of a blood clot. No additional information was provided. It is not known whether the autopsy was performed. Given alternative explanation and risk factors of pregnancy, labor and skeletal injury (trauma) the event of thrombosis is considered inconsistent with the causal association to immunization, per the WHO causality classification for adverse events following immunization. Events of skeletal injury was result of an accident and therefore not considered related. Company causality for event of thrombosis is considered not related to Janssen COVID-19 vaccine (Level 4 -Insufficient information available to confirm a possible, probable or a definitive case of venous thrombosis, per the Brighton Collaboration case definition); Reported Cause(s) of Death: BLOOD CLOTS; BROKE TAIL BONE

VAERS ID: 1227925 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Kansas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Platelet count, Pulmonary embolism, Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Systemic lupus erythematosus (broad), Embolic and thrombotic events, venous (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: General physical health deterioration
Preexisting Conditions: Medical History/Concurrent Conditions: Cancer (Unspecified type)
Allergies:
Diagnostic Lab Data: Test Name: Platelet count; Result Unstructured Data: Thrombocytopenia (count unspecified)
CDC Split Type: USJNJFOC20210427876

Write-up: SUSPECTED PULMONARY EMBOLISM; THROMBOCYTOPENIA; This spontaneous report received from a health care professional from a state immunization program concerned a 60 year old male. The patient"s height, and weight were not reported. The patient"s past medical history included cancer, and concurrent conditions included overall poor health. The patient received COVID-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown), date of administration was not reported, for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, approximately 24 hours post-vaccination, the patient was taken to the hospital after being found unresponsive. It was suspected that the patient had experienced a pulmonary embolism, resulting in his death. Laboratory results revealed thrombocytopenia. Reportedly, no heparin was used in his treatment. On an unspecified date, the subject died from suspected pulmonary embolism, and had not recovered from thrombocytopenia. An autopsy had not been performed at the time of the report, as it was pending family"s approval. The action taken with COVID-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: V0: This is a spontaneous report of a 60-year-old male patient, who died of a suspected pulmonary embolism approximately 24 hours after receipt of the Janssen COVID-19 vaccine. The patient had an unspecified cancer and was described as in overall poor health. He was brought to a hospital, where it was suspected that he had died due to a pulmonary embolism. He was also found to be thrombocytopenic. The patient"s cancer could provide a plausible alternative explanation for the event, although there are insufficient details to make a meaningful medical assessment at this time.; Reported Cause(s) of Death: SUSPECTED PULMONARY EMBOLISM

VAERS ID: 1227926 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-04-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210428310

Write-up: BLOOD CLOT; This spontaneous report received from a consumer via a company representative and concerned a male of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, a week or so after the Covid-19 vaccination the patient passed away in his sleep. The patient had no underlying condition. An autopsy was performed on an unspecified date and the patient was found to have blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: A male patient of unspecified age passed away in his sleep an unspecified time after receiving the Janssen COVID-19 vaccine for prevention of COVID-19 infection. It stated that the patient had no underlying condition. A blood clot was found by autopsy; no further details are provided. There is insufficient information to make a meaningful medical assessment. Additional information is being sought.; Reported Cause(s) of Death: BLOOD CLOT; Autopsy-determined Cause(s) of Death: BLOOD CLOT

VAERS ID: 1227927 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: North Carolina
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-04-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Caffeine consumption (heavy); Feeling unwell (upon waking the morning prior to receiving vaccination.); Heart disorder; Heavy smoker
Preexisting Conditions: Medical History/Concurrent Conditions: Stroke
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210428883

Write-up: PASSED AWAY; This spontaneous report was received from a partner, and concerned a 64 year old patient of an unspecified sex. The patient''s height, and weight were not reported. The patient''s past medical history included strokes; concurrent conditions included heavy smoker, caffeine consumption, unspecified heart issues, and felling unwell. The patient received Covid-19 vaccine Ad26.COV2.S (suspension for injection, route of admin, and batch number were not reported) dose, vaccination site, and start therapy date were not reported, for prophylactic vaccination. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient passed away. It was reported that the patient received Covid-19 vaccine in the morning and passed away later that day. The patient had felt unwell upon waking up that morning prior to vaccination. The coroner ruled out the vaccine as a potential cause of death due to the individual''s past and concurrent medical conditions. It was unspecified if an autopsy was performed. Action taken with Covid-19 vaccine Ad26.COV2.S was not applicable. This report was serious (Death).; Sender''s Comments: V0: A 64-year-old patient of unspecified sex received the Janssen COVID vaccine in the morning and died later that day. The coroner ruled out the vaccine as a potential cause of death due to the individual''s past and concurrent medical conditions, although these conditions weren''t specifically reported. Given the coroner''s assessment and symptoms preceding vaccination, there is a plausible alternate explanation for the death.; Reported Cause(s) of Death: PASSED AWAY

VAERS ID: 1227966 (history)
Form: Version 2.0
Age:

Sex: Male
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-19

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? 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:
CDC Split Type: USPFIZER INC2021404945

Write-up: got the shot for covid and died a few weeks later; This is a spontaneous report from a contactable consumer. A male patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. The relevant medical history and concomitant medications were reported as none. The patient got the shot for covid and died a few weeks later on an unspecified date. It was unknown if an autopsy was performed. The outcome of the event was fatal. Information on the Lot/Batch number has been requested.; Reported Cause(s) of Death: got the shot for covid and died a few weeks later

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VAERS ID: 1232815 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-04-20

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	1802068 / 1	LA / IM

Administered by: Private **Purchased by:** ?
Symptoms: Abdominal pain, Blood lactic acid, Blood phosphorus increased, Blood pressure immeasurable, Coma, Death, Headache, Hypotension, Shock, Syncope, Vomiting
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (narrow), Acute pancreatitis (broad), Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Retroperitoneal fibrosis (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Torsade de pointes, shock-associated conditions (broad), Hypovolaemic shock conditions (narrow), Toxic-septic shock conditions (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypoglycaemic and neurogenic shock conditions (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Chronic kidney disease (broad), Hypersensitivity (narrow), Tumour lysis syndrome (broad), Hypoglycaemia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: BP unobtainable, pH 6.8, lactate 15 Contact Hospital for lab and clinical details
CDC Split Type:

Write-up: Abdominal pain, vomiting, headache, hypotension, syncope, coma, shock, death at 2:00 pm 03/08/2021

VAERS ID: 1235825 (history)
Form: Version 2.0
Age:
Sex: Female
Location: New Hampshire
Vaccinated: 2021-03-26

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9269 / 1	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Bradycardia, Hypotension, Investigation
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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 Name: critical labs; Result Unstructured Data: Test Result:Unknown results
CDC Split Type: USPFIZER INC2021412017

Write-up: bradycardia; hypotension; This is a spontaneous report from a contactable nurse (Registered nurse with title of Infection Preventionist). A female patient of an unspecified age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL9269; expiration date: 01May2021) via an unspecified route of administration on 26Mar2021 at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient was with bradycardia, hypotension (seriousness criteria: hospitalization, death) and she passed away in the emergency room (ER), critical labs, she didn''t even make it one day, the reporter send her out and she passed away in the ER. The patient died on an unspecified date. It was not reported if an autopsy was performed.; Sender''s Comments: A causal relationship between BNT162B2 and the events hypotension,bradycardia cannot be excluded based on 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 RAs, Ethics Committees, and Investigators, as appropriate .; Reported Cause(s) of Death: bradycardia; hypotension

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VAERS ID: 1235830 (history)

Form: Version 2.0
Age:
Sex: Male
Location: South Carolina
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Drug ineffective, SARS-CoV-2 test, Suspected COVID-19
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Atrial fibrillation (Underlying health issues); Pre-diabetes (Underlying health issues)
Allergies:
Diagnostic Lab Data: Test Name: two weeks after testing positive for the virus; Test Result: Positive
CDC Split Type: USPFIZER INC2021415071

Write-up: died about a month ago; two weeks after testing positive for the virus; two weeks after testing positive for the virus; This is a spontaneous report from a non-contactable consumer. An 80-years-old male patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date at SINGLE DOSE for covid-19 immunisation. Medical history included pre-diabetes and atrial fibrillation (Underlying health issues). The patient"s concomitant medications were not reported. The patient previously received first dose of bnt162b2 on an unspecified date at SINGLE DOSE for covid-19 immunisation. The patient died about a month ago, two weeks after testing positive for the virus. The death occurred about four weeks after the second dose of the Pfizer/BNT vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Deceased

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VAERS ID: 1235832 (history)

Form: Version 2.0
Age:
Sex: Female
Location: New Hampshire
Vaccinated: 2021-03-26
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9269 / 1	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Confusional state, Dyspnoea, Hypoxia, Sepsis
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Hypoglycaemia (broad), Infective pneumonia (broad), Sepsis (narrow), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: USPFIZER INC2021415135

Write-up: wound up in the ER with hypoxia and sepsis; wound up in the ER with hypoxia and sepsis; shortness of breath; increased confusion; This is a spontaneous report from a contactable Nurse. A female patient of an unspecified age received first dose of bnt162b2 (BNT162B2), via an unspecified route of administration on 26Mar2021 (Lot Number: EL9269; Expiration Date: 01May2021) as single dose for Covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. It was reported that the female patient who we sent out with shortness of breath and increased confusion, she wound up in the ER (emergency room) with hypoxia and sepsis and she passed away. The events were serious as hospitalization and death. The patient died on an unspecified date. It was not reported if an autopsy was performed.; Sender''s Comments: Based on the limited available information, the Company considered there was not a

reasonable possibility that the reported events were related to the suspect product BNT162B2 (COMIRNATY). 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.; Reported Cause(s) of Death: increased confusion; shortness of breath; wound up in the ER with hypoxia and sepsis; wound up in the ER with hypoxia and sepsis

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VAERS ID: 1245485 (history)
Form: Version 2.0
Age:
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: death after 1 dose of vaccine; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (death after 1 dose of vaccine) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient''s past medical history included No adverse event. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (death after 1 dose of vaccine) in a

male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient''s past medical history included No adverse event. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Not Provided The reporter stated that she was not sure of the vaccine name that the patient used.; Sender''s Comments: This is a male patient of unknown age, who received mRNA-1273 Vaccine and died on an unknown date after receiving first dose of vaccine. No medical hx or conmeds were provided. The fatal outcome may be related to the patient''s pre-existing comorbidities Very limited information has been reported at this time. Further information is expected.; Reported Cause(s) of Death: death after 1 dose of vaccine

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VAERS ID: 1249228 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-23

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210426837

Write-up: DEATH; This spontaneous report received from a physician via a company representative concerned a female of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, once total, administered on 2021 for prophylactic vaccination. The batch number

was not reported and has been requested. No concomitant medications were reported. On an unspecified date, three days after vaccination, the patient died of unknown cause. It was unknown, whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1249255 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-23

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210428967

Write-up: DIED IN SLEEP; This spontaneous report received from a patient via a company representative concerned a female of unspecified age. The patient"s weight, height, and medical history were not reported. Patient had children. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: UNKNOWN, expiry: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Company is unable to perform follow-up to confirm batch/lot number. No concomitant medications were reported. On an unspecified date, the patient died in her sleep from an unknown cause of death. It was unknown if an

autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210429895.; Sender''s Comments: V0: 20210428967-COVID-19 VACCINE AD26.COV2.S-Died in sleep. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1249280 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-23

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

Administered by: Other **Purchased by:** ?
Symptoms: Coronary artery occlusion
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210431438

Write-up: HEART ATTACK FROM BLOOD CLOT; This spontaneous report received from a patient via a company representative concerned a Not Hispanic or Latino and Asian male of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient experienced heart attack from blood clot and the patient died. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s

was not applicable. This report was serious (Death).; Sender"s Comments: V0: This is an Asian male, unspecified age, who experienced a heart attack from a blood clot on an unspecified date after receiving the covid-19 vaccine ad26.cov2.s also on an unspecified date. No other details given. The information provided precludes a meaningful medical assessment. Additional information will be requested.; Reported Cause(s) of Death: DEATH

VAERS ID: 1249697 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Virginia
Vaccinated: 2021-04-09
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Intentional self-injury
SMQs:, Suicide/self-injury (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-04-18
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: passed away from self inflicted injury; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by an other health care professional (subsequently medically confirmed) and describes the occurrence of INTENTIONAL SELF-INJURY (passed away from self inflicted injury) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient"s past medical history included No adverse event. On 09-Apr-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced INTENTIONAL SELF-INJURY (passed away from self inflicted injury) (seriousness criterion death). The patient died on 18-Apr-2021. The cause of death was not reported. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter

did not provide any causality assessments. Concomitant product use was not provided by the reporter. No treatment information was provided. Action taken with mRNA-1273 in response to the event was not applicable. Reporter did not allow further contact; Sender''s Comments: Patient died of an unknown cause from a self-inflicted injury nine days after receiving second dose of Moderna Vaccine. Very limited information regarding this event has been provided at this time. Further information can not be requested.; Reported Cause(s) of Death: cause of death unknown

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VAERS ID: 1253991 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-24

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: DEATH; This spontaneous report received from a consumer concerned two patients unspecified age and sex. This report received via social media.. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown) dose, start therapy date were not reported, 1 total for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. It was reported that two people died recently after the vaccine. The cause of death was unknown. It was unspecified if an autopsy was performed The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report

was serious (Death).; Sender"s Comments: V0: -covid-19 vaccine ad26.cov2.s-death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1254024 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Completed suicide, Tinnitus
SMQs:, Suicide/self-injury (narrow), Hearing impairment (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: committed suicide; ringing in his ears; This spontaneous case was reported by a consumer and describes the occurrence of COMPLETED SUICIDE (committed suicide) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced COMPLETED SUICIDE (committed suicide) (seriousness criterion death) and TINNITUS (ringing in his ears). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, TINNITUS (ringing in his ears) outcome was unknown. Concomitant medications were

not reported; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1254028 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Hypotension, Thrombosis
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATXINC.MOD2021083

Write-up: Patient passed away; patient had blood clots in his brain/ Legs/ Lungs/ Arms; Severe hypotension; This spontaneous case was reported by a physician (subsequently medically confirmed) and describes the occurrence of DEATH (Patient passed away), THROMBOSIS (patient had blood clots in his brain/ Legs/ Lungs/ Arms) and HYPOTENSION (Severe hypotension) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient''s past medical history included No adverse event. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced DEATH (Patient passed away) (seriousness criterion death), THROMBOSIS (patient had blood clots in his brain/ Legs/ Lungs/ Arms) (seriousness criterion death) and HYPOTENSION (Severe hypotension) (seriousness criterion death). The cause of death was not reported. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality

assessments. Treatment information was not provided. Company comment: This case concerns the death of a patient of unknown age and gender after administration of mrna-1273 ()LOT UNKNOWN). Very limited information regarding this event/s has been provided at this time. Further information has been requested. Critical details such as the mRNA-1273 date of administration, onset of any signs an d symptoms, and date of death is lacking.; Sender''s Comments: This case concerns the death of a patient of unknown age and gender after administration of mrna-1273 ()LOT UNKNOWN) . Very limited information regarding this event/s has been provided at this time. Further information has been requested. Critical details such as the mRNA-1273 date of administration, onset of any signs an d symptoms, and date of death is lacking.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1255809 (history)
Form: Version 2.0
Age: 62.0
Sex: Female
Location: Wisconsin
Vaccinated: 2021-04-16
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-24

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EW0158 / 2	LA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Death, Fatigue, Headache, 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: 2021-04-22
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies: simvastatin
Diagnostic Lab Data:
CDC Split Type:

Write-up: -Patient died unexpectedly on Thursday, April 22, 2021 at her home (found unresponsive by spouse) -No reactions were observed at the pharmacy (within 15 min waiting period) following either covid-19 vaccine (1st dose administered 8am on 3/26/21 -Left Deltoid) -When asked (4-16-21) how she tolerated the 1st dose, Patient reported minor side effects (mild Headache, tiredness) but had no reservations about receiving the 2nd dose. No additional information.

VAERS ID: 1255617 (history)
Form: Version 2.0
Age:
Sex: Female
Location: New Hampshire
Vaccinated: 2021-03-26
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9269 / 1	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Cardiac failure congestive, Hypotension, Hypoxia
SMQs:, Cardiac failure (narrow), Anaphylactic reaction (broad), Asthma/bronchospasm (broad), Neuroleptic malignant syndrome (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Respiratory failure (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: she ended up passing away in the hospital, her diagnosis there was hypoxia and acute CHF; she ended up passing away in the hospital, her diagnosis there was hypoxia and acute CHF; hypotensive; This is a spontaneous report from a contactable Nurse. A female

patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 26Mar2021 (Lot Number: EL9269; Expiration Date: 01May2021) as single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient sent to the ER hypoxic, hypotensive, short of breath, she ended up passing away in the hospital, her diagnosis there was hypoxia and acute CHF (Congestive heart failure). Serious: No. The patient died on an unspecified date. It was not reported if an autopsy was performed.; Sender''s Comments: Based on the information available, a causal association between BNT162B2 and the reported events cannot be excluded. However, details on the patent''s age, medical history, drug-event temporal relationship, clinical course of the event and relevant test results would allow for a meaningful 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 RAs, Ethics Committees, and Investigators, as appropriate.; Reported Cause(s) of Death: she ended up passing away in the hospital, her diagnosis there was hypoxia and acute CHF; she ended up passing away in the hospital, her diagnosis there was hypoxia and acute CHF

VAERS ID: 1255618 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New Hampshire
Vaccinated: 2021-03-26
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9269 / 1	- / -

Administered by: Senior Living **Purchased by:** ?
Symptoms: Cerebral artery occlusion, Cerebrovascular accident
SMQs:, Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: I have another male (patient) who had an acute CVA we send him to the hospital he had acute CVA, he had a right artery occlusion, he passed away; I have another male (patient) who had an acute CVA we send him to the hospital he had acute CVA, he had a right artery occlusion, he passed away; This is a spontaneous report from a contactable Nurse (Registered nurse with title of Infection Preventionist). A male patient of an unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 26Mar2021 (Lot Number: EL9269; Expiration Date: 01May2021) as single dose for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. The patient had an acute CVA and was sent him to the hospital he had acute Cerebrovascular accident (CVA), he had a right artery occlusion, he passed away. The patient died on an unspecified date in 2021. It was not reported if an autopsy was performed.; Sender"s Comments: The information currently available is very limited. There is no sufficient evidence that the reported events may be related to administration of BNT162B2. Of note, medical history and concomitant medications were not provided to determine pre-existing risk factors or conditions that may have led to the events. Case will be re-assess 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 RAs, Ethics Committees, and Investigators, as appropriate.; Reported Cause(s) of Death: patient had an acute CVA and was sent him to the hospital he had acute CVA, he had a right artery occlusion; patient had an acute CVA and was sent him to the hospital he had acute CVA, he had a right artery occlusion

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VAERS ID: 1255703 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Portal vein thrombosis, Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Systemic lupus erythematosus (broad), Embolic and thrombotic events, venous (narrow), Drug reaction with eosinophilia and

systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021425367

Write-up: Patient died; portal vein thrombosis; thrombocytopenia; This is a spontaneous report from a contactable physician. A 50-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on an unspecified date as SINGLE DOSE for covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. The patient experienced portal vein thrombosis and thrombocytopenia 2 weeks after first Pfizer vaccine. Patient died. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the events portal vein thrombosis and thrombocytopenia was unknown. The outcome of the event unknown cause of death was fatal. Information on the lot/ batch number has been requested.; Sender''s Comments: The information available in this report is limited and does not allow a medically meaningful assessment of the case. Based on temporal association, a causal association between the reported events and BNT162B2 cannot be fully excluded. Case will be reassessed when additional information is available including medical history and concomitant drug 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 RAs, Ethics Committees, and investigators, as appropriate. ; Reported Cause(s) of Death: Patient died

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VAERS ID: 1255705 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot /	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021426236

Write-up: died after taking the vaccine; This is a spontaneous report from a contactable consumer reporting on behalf of the mother. A female patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date, at single dose, for COVID-19 immunisation. Medical history and concomitant medications were not reported. The patient died after taking the vaccine, on an unspecified date. It was unknown if an autopsy was performed. Cause of death was unknown. The reporter believed that it was from the vaccine and ICU nurse said, doctor questioned about what the patient got sick and reporter just wanted to report it. Information about lot/batch number has been requested.; Reported Cause(s) of Death: died after taking the vaccine

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VAERS ID: 1255731 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?

Symptoms: COVID-19, Drug ineffective, Organ failure, Thrombosis
SMQs:, Lack of efficacy/effect (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), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021432758

Write-up: Hot a fever then full blown covid; Hot a fever then full blown covid; blood clot; organ failure; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that a male patient of an unspecified age received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. Previously on an unknown date, the patient received the first dose of BNT162B2 vaccine. On an unspecified date, the patient experienced hot a fever then full blown COVID, blood clot and organ failure leading to patient death on an unknown date. It was not reported if an autopsy was performed. No follow-up attempts are possible. Information about lot/batch number cannot be obtained. No further information is expected; Reported Cause(s) of Death: Drug ineffective; Covid-19; Blood clot; Organ failure

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VAERS ID: 1255738 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 2021-03-30
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / UNK	- / -
UNK: VACCINE NOT SPECIFIED (NO BRAND NAME) / UNKNOWN MANUFACTURER	2101912 / UNK	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Dyspnoea, Illness, Myalgia, Nausea, Pain, Vomiting
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (broad), Acute pancreatitis (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Cardiomyopathy (broad), Eosinophilic pneumonia (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: ADEMPAS; AMLODIPINE BESILATE; XARELTO
Current Illness: Pulmonary arterial hypertension
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021434134

Write-up: Nausea; Vomiting; Shortness of breath; Muscle aches; Extremely ill; Body aches; expired; This is a report from a contactable consumer based on the information received by Pfizer. (Manufacturer Report Number: UNT-2021-006444). A 76-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for Covid-19 immunization; and treprostinil sodium (TYVASO, strength: 0.6, mg/ml), via resp inhalation (reported as via inhalation route) from 30Mar2021 (Batch/Lot Number: 2101912; Expiration Date: 31Jan2022) to an unspecified date, at 18-54 ug, four times a day (QID) for primary pulmonary arterial hypertension. Medical history included ongoing pulmonary arterial hypertension. Concomitant medications included riociguat (ADEMPAS), amlodipine besilate, rivaroxaban (XARELTO); all taken for an unspecified indication, start and stop date were not reported. It was reported that the patient began therapy with IH Tyvaso (treprostinil sodium, concentration of 0.6 mg/ml) delivered by Tyvaso Inhalation Device (TD-300/A), on 30Mar2021 for primary pulmonary arterial hypertension. The current dose was reported as 18-54 ug (3-9 breaths), four times a day (QID) via inhalation (IH) route. On an unspecified date, the patient experienced nausea, vomiting, shortness of breath, muscle aches, extremely ill, and body aches. The patient was hospitalized in response to the events on an unspecified date. The outcome of events was unknown. On an unspecified date in 2021, the patient expired, and cause of death was not reported. The action taken in response to the events for treprostinil sodium was unknown. The patient died on an unspecified date in 2021. It was unknown if an autopsy was performed. The reporter''s assessment of the causal relationship between the events with the suspect product was not provided at the time of this report. Since no determination has been received, the case is managed based on the company causality

assessment. Case Comment/Senders Comment: The company has assessed the serious adverse event of death as not related to IH treprostinil and TD-300/A device. The event was likely due to progressive complications and life limiting nature of underlying PAH in this elderly patient. Information about lot/batch number has been requested.; Sender''s Comments: The information currently available is limited and does not allow a meaningful causality assessment for reported event of death (unknown cause); however, based solely on implied vaccine-event chronological association a causal relationship between this event and BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) cannot be completely excluded. The other reported events; Reported Cause(s) of Death: expired

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VAERS ID: 1255744 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021438418

Write-up: Deceased; This is a spontaneous report from a Pfizer. A non-contactable consumer reported that a patient of unspecified age and gender received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient previously received bnt162b2

(BNT162B2), dose 1 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunization. Consumer reported that, "Amongst my coworkers about 50% were put down for two to three days and one person became deceased a few hours after the second shot." The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow up attempts are possible; Information on the lot/batch number cannot be obtained.; Reported Cause(s) of Death: Deceased

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VAERS ID: 1255749 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021445170

Write-up: died after the second one when they had covid; died after the second one when they had covid; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), second dose via an unspecified route of administration on an unspecified date (batch/lot number was not reported) as a single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient died after the second one

when they had Covid because there were too many antibodies on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: died after the second one when they had covid; died after the second one when they had covid

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VAERS ID: 1261201 (history)
Form: Version 2.0
Age: 36.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-27

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210428954

Write-up: DEATH; This spontaneous report received from a patient concerned a 36 year old male. The patient''s weight, height, and medical history were not reported. As per reporter patient was healthy. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported, per procedure no follow up will be requested for this case. No concomitant medications were reported. The reported stated that presence of antibodies should be tested before taking the vaccine and taking the vaccine with covid antibodies present can kill. The patient died on an unspecified date with unknown cause of death after vaccination. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210428954 -covid-19 vaccine ad26.cov2.s -Death. This event(s) is considered unassessable. The event(s) has a

compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1261290 (history)
Form: Version 2.0
Age: 63.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-27

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210436493

Write-up: DEATH; This spontaneous report received from a patient concerned a 63 year old female. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot number. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death. It was reported that, the patient death occurred after receiving the vaccine. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked.; Sender"s Comments: V0: -covid-19 vaccine ad26.cov2.s-death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the

event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1261835 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-27

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Injury
SMQs:, Accidents and injuries (narrow), Hostility/aggression (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021448793

Write-up: who is responsible and/or liable when a child is injured or dies during a COVID vaccine trial; who is responsible and/or liable when a child is injured or dies during a COVID vaccine trial; This is a spontaneous report from a contactable consumer or other non hcp. A child patient of an unspecified gender received bnt162b2 (BNT162B2), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. It was reported that the consumer had searched the website and cannot find anywhere that states who is responsible and/or liable when a child is injured or dies during a COVID vaccine trial. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: who is responsible and/or liable when a child is injured or dies during a COVID vaccine trial

VAERS ID: 1266077 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021458298

Write-up: Had both vaccines and died from COVID-19 afterwards; Had both vaccines and died from COVID-19 afterwards; This is a spontaneous report from a contactable nurse. A 42-year-old patient of an unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 1 and dose 2; both via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient had both vaccines and died from COVID-19 afterwards on an unspecified date. The patient did not have any known underlying health conditions or issues. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/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. This 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 to regulatory authorities, Ethics Committees, and Investigators, as appropriate.; Reported Cause(s) of Death: Had both vaccines and died from COVID-19 afterwards; Had both vaccines and died from COVID-19 afterwards

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VAERS ID: 1266078 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Arkansas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Dyspnoea, Pulmonary embolism
SMQs:, Anaphylactic reaction (broad), Embolic and thrombotic events, venous (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021459420

Write-up: pulmonary embolism; short of breath; This is a spontaneous report from a non-contactable consumer via Pfizer sales representative. A 63-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient died of pulmonary embolism 2 days after 2nd vaccine dose. She had been short of breath the day after the shot, called the pharmacy where she"d

received it, and they told her to go to ER. The reporter thought that she did not go to the ER, and then later died at home. It was unknown if an autopsy was performed. Outcome of the event short of breath at the time of death was unknown. No follow-up attempts are possible; Information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: pulmonary embolism

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VAERS ID: 1269775 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-29

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021465549

Write-up: dies hours after getting Covid-19 vaccine; This is a spontaneous report from a non-contactable consumer (Pfizer-sponsored program). A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; Solution for injection; Lot Number: Unknown), via an unspecified route of administration on an unspecified date as SINGLE DOSE for covid-19 immunization. The patient medical history and concomitant medications were not reported. The patient experienced dies hours after getting covid-19 vaccine on an unspecified date. It was not reported if an autopsy was performed. The outcome of the event was fatal. No follow-up attempts are Possible; information about batch/ lot number cannot be obtained.; Reported Cause(s) of Death: dies hours after getting Covid-

19 vaccine

VAERS ID: 1269944 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Arizona
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210444879

Write-up: PASSED AWAY/ DEATH; This spontaneous report received from a consumer via company representative concerned a 70 year old female. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry date: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot number. No concomitant medications were reported. On an unspecified date, the patient passed away after receiving the vaccine. The patient died from an unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0:-covid-19 vaccine ad26.cov2.s-death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1269961 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210450529

Write-up: DEATH; This spontaneous report received from a pharmacist concerned a male of an unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported,1 total, administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient who received the JANSSEN COVID-19 vaccine and passed away a few weeks later. It was not sure whether his other medical issues contributed to his death. The patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: - COVID-19 VACCINE AD26.COV2.S - Death, This event is considered Unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1272739 (history)

Form: Version 2.0
Age: 77.0
Sex: Female
Location: Florida
Vaccinated: 2021-04-28
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-04-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	048A21A / 1	UN / SYR
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	001C21A / 2	UN / SYR

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-04-29
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Symbicort Allopurinol, Pantoprazole, Furosemide,
Current Illness:
Preexisting Conditions: COPD, HTN, DM, A-FIB
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Expired within 72 hours of receiving vaccine.

VAERS ID: 1276412 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Alabama
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

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

Symptoms: Death, Hospitalisation
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210443181

Write-up: HOSPITALIZATION; DEATH; This spontaneous report received from a patient via a company representative concerned a female of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, 01 total, for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. After receiving the vaccine, on an unspecified date, the patient was hospitalized. The patient died after 6 days of hospitalization. The reason of death was unknown. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome was fatal. This report was serious (Death, and Hospitalization Caused / Prolonged).; Sender''s Comments: V0:-JANSSEN COVID-19 VACCINE Ad26.COV2.S- Death, Hospitalization. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1276505 (history)
Form: Version 2.0
Age: 60.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

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

Symptoms: Myocardial infarction
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210452359

Write-up: HEART ATTACK; This spontaneous report (social media) received from a patient via a company representative concerned a 60 year old male. The patients weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported batch number: Unknown) dose was not reported, 1 total administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient took vaccine and at night patient experienced heart attack and died due to heart attack. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to; Sender''s Comments: V0-Covid-19 vaccine ad26.cov2.s -Heart attack. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: HEART ATTACK

VAERS ID: 1276508 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Adverse reaction, Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210452419

Write-up: DEATH; UNSPECIFIED COMPLICATIONS; This spontaneous report received from a consumer concerned multiple patients. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, once total, for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. The six patients died on unspecified date due to unknown cause. It was also stated that these patients had some unspecified complications. It was unknown, whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patients died of death on an unspecified date, and the outcome of complications was not reported. This report was serious (Death).; Sender"s Comments: V0: 20210452419-covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1276513 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Malaise

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210452604

Write-up: DEATH; FELT SICK; This spontaneous report received from a consumer (social media) via a company representative concerned a male of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, Expiry date: Unknown) frequency 1 total, dose and start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, after getting the vaccine, the patient felt sick. On the next dat, the patient was deceased. The cause of death was not reported. It was unknown whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: -covid-19 vaccine ad26.cov2.s -Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1276522 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Adverse reaction, Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210453204

Write-up: DEATH; COMPLICATIONS FROM THE VACCINE; This spontaneous report received from a consumer concerned an adult male. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, once total for prophylactic vaccination. The batch number was not reported. In this scenario follow-up is not required to obtain batch/lot numbers. No concomitant medications were reported. On an unspecified date, two months after vaccination, the patient died of unknown cause. The patient death occurred in hospital. It was also stated that patient had some unspecified complications. iT was also reported that, patient was healthy man in his early sixties. It was unknown, whether autopsy was performed or not. On an unspecified date, the subject experienced death, and complications from the vaccine. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of death on an unspecified date, and the outcome of complications from the vaccine was not reported. This report was serious (Death).; Sender''s Comments: V0:-covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s); Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1276530 (history)
Form: Version 2.0
Age: 40.0
Sex: Female
Location: Oregon
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210453777

Write-up: DEATH; BLOOD CLOT; This spontaneous report received from a consumer news/social media platform concerned a 5 decade old female. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient developed a rare blood clot and died within two weeks of getting the Janssen covid vaccine. On an unspecified date, the patient died from unknown cause of death. It was unknown whether autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Death, and Other Medically Important Condition). This case, from the same reporter is linked to.; Sender"s Comments: V0-covid-19 vaccine ad26.cov2.s-This case concerns with 5 decade old female. Death, Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1276535 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Unknown cause of death
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210453979

Write-up: DEATH; This spontaneous report received from a consumer via social media concerned a female of unspecified age. The patient"s height, and weight were not reported. The patient"s past medical history included unknown cause of death. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: UNKNOWN) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. Consumer posted on social media that the J and J vaccine killed her mother and they are waiting on her autopsy . The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0- 20210453979 - Covid-19 vaccine ad26.cov2.s-death.This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.

VAERS ID: 1276582 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: unknown cause of death.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210456256

Write-up: DEATH; This spontaneous report received from a consumer concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. The patient''s pre-existing medical conditions included unknown cause of death. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient received the Janssen COVID-19 vaccine and then died 15 days later. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). .; Sender''s Comments: V0-20210456256- COVID-19 VACCINE AD26.COV2.s.adverse effect . This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1276796 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and

venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: blood clot; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clot) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (blood clot) (seriousness criterion death). The reported cause of death was Clot blood. It is unknown if an autopsy was performed. The reporter stated that his wife had recently passed away from a blood clot after receiving a second dose of the Moderna COVID-19 vaccine. Treatment information was not provided. Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Clot blood

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VAERS ID: 1279433 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-02

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Heart rate, Heart rate increased

SMQs:, Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: heartbeat; Result Unstructured Data: Test Result:racing
CDC Split Type: USPFIZER INC2021467483

Write-up: racing heartbeat; This is a spontaneous report from a contactable consumer via Medical information team. A female patient of an unspecified age (reported as age: 55, unit: unknown) received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient died a few days later after getting vaccinated from a racing heartbeat. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information about batch/lot number has been requested.; Reported Cause(s) of Death: racing heartbeat

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VAERS ID: 1279434 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Utah
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-02

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021467484

Write-up: Died after the 2nd dose; This is a spontaneous report from a contactable consumer from a Pfizer-sponsored program. A 39-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient previously received the first dose of BNT162B2 on an unspecified date for COVID-19 immunization. The patient died after the 2nd dose on an unspecified date. The cause of death was not reported. It was not reported if an autopsy was performed. Information on the batch/lot number has been requested.; Reported Cause(s) of Death: died after the 2nd dose

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VAERS ID: 1285078 (history)
Form: Version 2.0
Age: 92.0
Sex: Male
Location: California
Vaccinated: 2021-01-11
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-03

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 1	RA / IM
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	RA / IM

Administered by: Other **Purchased by:** ?
Symptoms: Atrial flutter, Death, Glucose tolerance impaired, Haemorrhage intracranial, Pain in extremity, Peripheral artery thrombosis, Peripheral coldness, Thrombectomy, Thrombophlebitis superficial, Ultrasound Doppler abnormal, Vasodilatation
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Hyperglycaemia/new onset diabetes mellitus (narrow), Supraventricular tachyarrhythmias (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and

venous (narrow), Embolic and thrombotic events, venous (narrow), Cardiomyopathy (broad), Lipodystrophy (broad), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: propranolol, other antihypertensive, antacid for GERD
Current Illness: none
Preexisting Conditions: hypertension, GERD, borderline DM, diagnosed while in rehab week before death - well controlled/no Rx.
Allergies: none
Diagnostic Lab Data: Available from Dr.
CDC Split Type:

Write-up: Patient was a well and active 92 year old man, who walked 2 miles daily with a walker. He received Covid vaccine dose 1 on 11 Jan. On or before 30 Jan, he noted pains in one leg while walking. Newly dilated veins on the lower leg were noted. On feb 2 a doppler US revealed a superficial saphenous vein venous thrombosis, also new onset A flutter. Aspirin begun. On Feb 5 he woke with a cold foot (same side) and was found to have an arterial thrombus from SFA through distal posterial tibial. He underwent emergency thrombectomy, went to rehab for recovery, on Feb 23 suffered intracranial hemorrhage and died on Mar 10.

VAERS ID: 1284660 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-04

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210456400

Write-up: CLOTS; This spontaneous report received from a physician concerned a patient of unspecified age and sex. The patient"s weight, height and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown) frequency one total, dose, start therapy date were not reported for prophylactic vaccination.The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient died due to clot. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died to clots on an unspecified date. This report was serious (Death). This case, from the same reporter is linked to 20210457363.; Sender"s Comments: V0: 20210456400-Covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: CLOTS

VAERS ID: 1284730 (history)
Form: Version 2.0
Age: 68.0
Sex: Male
Location: Florida
Vaccinated: 2021-03-18
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Death; This case was received via an unknown source (no reference has been entered for a health authority or license partner) on 28-Apr-2021 and was forwarded to Moderna on 28-Apr-2021. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death) in a 68-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On 18-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No treatment information was provided. No concomitant medication was provided.; Sender''s Comments: This is a 68-year-old, male patient who received mRNA-1273 vaccine (batch no. unknown) who died, on an unknown date after receiving first dose of vaccine. No medical history or conmeds were provided. Existing comorbidities probably could have been the causative factor in his death. Very limited information has been reported at this time. Further information is expected.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1284731 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: General physical health deterioration, Product dose omission issue
SMQs:, Medication errors (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: General physical health deterioration (It is reported that patient had history of many medical conditions, unspecified.); Rehabilitation therapy (Patient was in rehabilitation center)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Died due to his many medical conditions; could not administer second vaccine; This spontaneous case was reported by a pharmacist (subsequently medically confirmed) and describes the occurrence of GENERAL PHYSICAL HEALTH DETERIORATION (Died due to his many medical conditions) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient"s past medical history included Rehabilitation therapy (Patient was in rehabilitation center) and General physical health deterioration (It is reported that patient had history of many medical conditions, unspecified.). On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced GENERAL PHYSICAL HEALTH DETERIORATION (Died due to his many medical conditions) (seriousness criterion death) and PRODUCT DOSE OMISSION ISSUE (could not administer second vaccine). The patient died on an unknown date. The reported cause of death was medical conditions. It is unknown if an autopsy was performed. At the time of death, PRODUCT DOSE OMISSION ISSUE (could not administer second vaccine) had resolved. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant product use was not provided by the reporter. No treatment information was provided.; Sender"s Comments: This is a male patient of unknown age who received mRNA-1273 vaccine (batch no. unknown) who died on an unknown date after receiving first dose of vaccine. Patient had history of many medical conditions. No conmeds were given. Reporter could not administer second vaccine. Very limited information has been reported at this time. No further information is expected.; Reported Cause(s) of Death: medical conditions

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VAERS ID: 1284865 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Tennessee
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-04

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

Administered by: Unknown **Purchased by:** ?

Symptoms: Haemorrhagic stroke
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhagic 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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: PLAVIX
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: CVA
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021477606

Write-up: hemorrhage stroke; This is a spontaneous report from a Pfizer-sponsored program, received from a contactable nurse. A 55-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not provided), via an unspecified route of administration on an unspecified date at single dose for covid-19 immunisation. The patient''s medical history included CVA. The patient''s concomitant medication included oral clopidogrel bisulfate (PLAVIX) for CVA at 25 mg, 1x/day (25 mg once everyday). It was reported that a patient who had the Covid vaccine, 3 days later the patient had hemorrhage and she passed away. It was clarified that patient had a hemorrhage stroke and passed away. The reporter did not know the date the patient passed away, but this was called on the office on 16Apr2021. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Sender''s Comments: Based on the temporal relationship, a causal association between the reported hemorrhagic stroke and the suspect drug, BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), cannot be completely excluded. Of note, patient has a medical history of CVA. 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.; Reported Cause(s) of Death: hemorrhage stroke

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VAERS ID: 1284940 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-05-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: girl died after getting second dose of Moderna vaccine it was thought to be fault of Tylenol.; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (girl died after getting second dose of Moderna vaccine it was thought to be fault of Tylenol.) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant and treatment information was provided. Company comment:This is an age unknown, female patient who received mRNA-1273 vaccine (batch no. unk) who died, after receiving second dose of vaccine. No medical history and conmeds were provided. Existing comorbidities probably could have been the causative factor in her death. Very limited information has been reported at this time. Further information is expected.; Sender"s Comments: This is an age unknown, female patient who received mRNA-1273 vaccine (batch no. unk) who died, after receiving second dose of vaccine. No medical history and conmeds were provided. Existing comorbidities probably could have been the causative factor in her death. Very limited information has been reported at this time. Further information is expected.; Reported Cause(s) of Death: Reporter mentioned that a girl died after getting second dose of Moderna vaccine.that it was thought to be fault of Tylenol.

VAERS ID: 1287928 (history)
Form: Version 2.0
Age:

Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-05

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210461305

Write-up: DEATH; This spontaneous report received from Pfizer via social media from a consumer concerned a female of unspecified age. The patient''s height, and weight were not reported. The patient''s past medical history included unknown cause of death. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, administered on 22-MAR-2021 for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died. It was unknown if the autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Other Medically Important Condition).; Sender''s Comments: V0- 20210461305-Covid-19 vaccine ad26.cov2.s-death.This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1291614 (history)
Form: Version 2.0
Age:
Sex: Unknown

Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-06

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: had a friend who dropped dead after getting the vaccine; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (had a friend who dropped dead after getting the vaccine) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product use was not provided by the reporter. Action taken with mRNA-1273 in response to the event was not applicable. This is a patient of unknown age and gender who received mRNA-1273 vaccine (batch no. unk) who died on an unknown date after receiving the first dose of vaccine No medical hx or concomitant products Very limited information has been reported at this time. Further information is being followed up; Sender"s Comments: This is a patient of unknown age and gender who received mRNA-1273 vaccine (batch no. unk) who died on an unknown date after receiving the first dose of vaccine No medical hx or concomitant products Very limited information has been reported at this time. Further information is being followed up; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1294841 (history)
Form: Version 2.0
Age:

Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Death; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product use was not provided by the reporter. The reporter informed there were 2 deaths of those who had received their second dose of Moderna vaccine, same day, same location who died 2 hours after vaccine administration. Treatment information was not provided. This is a patient who received mRNA-1273 (COVID 19 Vaccine Moderna) (batch no. unk) and died after he received the second dose of vaccine. No Medical History information was reported. Concomitant product use was not provided by the reporter Very limited information has been reported at this time. Further information is expected.; Sender"s Comments: This is a patient who received mRNA-1273 (COVID 19 Vaccine Moderna) (batch no. unk) and died after he received the second dose of vaccine. No Medical History information was reported. Concomitant product use was not provided by the reporter Very limited information has been reported at this time. Further information is expected.; Reported Cause(s) of Death: Unknown cause of death

VAERS ID: 1298832 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-08

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210507041

Write-up: DEATH; This spontaneous report received from a company representative concerned a patient of unspecified age and sex. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry date: Unknown) dose, 1 total administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died after getting Janssen vaccine. The patient died from an unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210507041-covid-19 vaccine ad26.cov2.s-Death. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1299367 (history)

Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Malaise
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021497007

Write-up: received his second dose of the vaccine and then died; got very ill; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. The patient had unspecified underlying health issues. The patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunization and experienced COVID-19. The reporter"s mom"s cousin who was in his late 70s/ early 80s, had some underlying health issues. He received his second dose of the vaccine, got very ill, and died. Reporter was unsure if he received Pfizer vaccine or different brand. The outcome of got very ill was unknown. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: received his second dose of the vaccine and then died

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VAERS ID: 1299369 (history)

Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-05-02
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021498045

Write-up: got infected with covid and died yesterday; Looks like the vaccine may not be as effective with the new variants.; This is a spontaneous report from a non-contactable consumer. A male patient (infectious disease doctor) received two doses of BNT162B2 (Solution for injection, lot number: unknown, Expiration date: Unknown) on unknown dates . The patient travelled and got infected with covid and died yesterday (02May2021). The reporter stated that it looked like the vaccine might not be as effective with the new variants. After two Pfizer vaccines shots , this doctor travelled to see his parents and succumbed to the new variant. Outcome of the event was fatal. No follow up attempts are possible; Information about Lot and batch number could not be obtained. No further information is expected.; Reported Cause(s) of Death: got infected with covid and died yesterday.

VAERS ID: 1301967 (history)
Form: Version 2.0
Age:

Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-10

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

Administered by: Other **Purchased by:** ?
Symptoms: Intracardiac thrombus
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210506615

Write-up: BLOOD CLOT IN HEART; This spontaneous report received from a consumer concerned a female of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, administered on 2021 for prophylactic vaccination. The batch number was not reported and it has been requested. No concomitant medications were reported. On 2021, the patient died due to blood clot in heart. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0 20210506615-COVID-19 VACCINE AD26.COV2.S- blood clot in heart. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT IN HEART

VAERS ID: 1304956 (history)
Form: Version 2.0
Age: 42.0
Sex: Male

Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-11

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210510772

Write-up: BLOOD CLOT; This spontaneous report received from a company representative concerned a 42 year old male. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry were unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, between three to ten days later after vaccination the patient died due to blood clots. It was unknown, whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0;20210510772-covid-19 vaccine ad26.cov2.s-Thrombosis. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1305027 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Texas

Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-11

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: died after receiving the Moderna Covid-19 vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (died after receiving the Moderna Covid-19 vaccine) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Not Provided Concomitant medications were not reported. No treatment information was provided. Action taken with mRNA-1273 in response to the drug was not applicable. Company comment: This is a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. died on an unknown date after the first dose of vaccine. No Medical History or Concomitant medications were reported. Very limited information has been reported at this time. Further information is not expected. This case was linked to US-MODERNATX, INC.-MOD-2021-012533, US-MODERNATX, INC.-MOD-2021-020196, MOD21-10588, MOD21-086364, MOD21-086368 (Linked Report).; Sender''s Comments: This is a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. died on an unknown date after the first dose of vaccine. No Medical History or Concomitant medications were reported. Very limited information has been reported at this time. Further information is not expected. US-MODERNATX, INC.-MOD-2021-012533: US-MODERNATX, INC.-MOD-2021-020196: MOD21-10588: MOD21-086364: MOD21-086368:; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1307002 (history)

Form: Version 2.0
Age: 71.0
Sex: Female
Location: Missouri
Vaccinated: 2021-01-13
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9264 / 2	- / IM

Administered by: Unknown **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? Yes
Date died: 2021-05-04
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: No adverse reaction to vaccine. Pt positive for covid in april 2021. recieved 2 doses of pfizer vaccine (1/2021&2/2021)

VAERS ID: 1309191 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-12

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210506934

Write-up: PASSED AWAY; This spontaneous report received from a consumer via social media concerned a female patient of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration was not reported, batch number and expiry were unknown) dose, start therapy date were not reported, 1 total, administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date (a couple of days after vaccination), the patient passed away. The cause of death was unknown. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210508328.; Sender''s Comments: V0: 20210506934 -COVID-19 VACCINE AD26.COV2.S - Passed Away. This event is considered Unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1309628 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-12

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

PFIZER/BIONTECH		

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021498707

Write-up: died 3 weeks later after getting the vaccine.; This is a spontaneous report from a Pfizer-sponsored program. A contactable nurse reported that a patient (unknown age and gender) received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number unknown) at single dose for COVID-19 immunisation on unknown date. Relevant history and concomitant drugs were unknown. The patient died 3 weeks later after getting the vaccine. It was unknown if autopsy was performed or not.; Sender"s Comments: The causal relationship between BNT162B2 and the fatal event cannot be excluded as 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.; Reported Cause(s) of Death: died 3 weeks later after getting the vaccine.

VAERS ID: 1309632 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-12

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021501749

Write-up: patient got infected with covid; patient got infected with covid; This is spontaneous report from a non-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 dates (Batch/Lot number was not reported) at 1st dose, single and 2nd dose, single for covid-19 immunization. The patient"s medical history was not reported. There were no concomitant medications. While visiting abroad, patient got infected with covid and died on an unspecified date. Apparently, the patient received two Pfizer vaccines shots prior to his travel. It was unknown if an autopsy was performed. No follow-up attempts are possible, information about batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: patient got infected with covid

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VAERS ID: 1312771 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-13

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Thrombosis

SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210510294

Write-up: FEMALE ISSUES; BLOOD CLOTS; This spontaneous report received from a consumer concerned multiple patients. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported and has been requested. On an unspecified date, the consumer called and reported that he had read in newspaper that women had gotten blood clots and 3 passed away with female issues. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of female issues on an unspecified date, and the outcome of blood clots was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: V0. 20210510294 -COVID-19 VACCINE AD26.COV2.S- Female issues, Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1314475 (history)
Form: Version 2.0
Age: 95.0
Sex: Female
Location: Puerto Rico
Vaccinated: 2021-02-03
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-13

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

Administered by: Senior Living **Purchased by:** ?

Symptoms: Death, SARS-CoV-2 test negative
SMQs:, COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-28
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Carvedilol, losartan 12.5mg, nitroglycerine, Atrovent, indur.
Current Illness: Unknown.
Preexisting Conditions: Congestive cardiac arrest.
Allergies: Unknown.
Diagnostic Lab Data: Covid-19 antigent test. Negative result.
CDC Split Type:

Write-up: Unknown

VAERS ID: 1315725 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-13

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210517449

Write-up: PASSED AWAY BECAUSE OF CLOT; This spontaneous report received from a consumer concerned a patient of Unspecified Race, ethnic origin, age and sex. The patient"s weight, height and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry: unknown) dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported and has been requested. On an unspecified date, the patient passed away because of clot. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210517449-covid-19 vaccine ad26.cov2.s-thrombosis. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: CLOT

VAERS ID: 1316349 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-14

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210514012

Write-up: This spontaneous report received from a consumer via a company representative concerned a female of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: Unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died from blood clot. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0. 20210514012-COVID-19 VACCINE AD26.COV2.S-Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1316353 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-14

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:

Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210516437

Write-up: DIED AFTER COMPLICATION FROM THE JOHNSON AND JOHNSON COVID-19 VACCINE; This spontaneous report received from a patient via a company representative concerned a female of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered on 2021 for prophylactic vaccination. No concomitant medications were reported. On 2021, the patient died after complication from the Johnson & Johnson COVID-19 vaccine. The cause of death was unknown. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0;20210516437-covid-19 vaccine ad26.cov2.-Died after complication from the Johnson & Johnson COVID-19 vaccine. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: DIED

VAERS ID: 1319776 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-15

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident, Death, Extra dose administered
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), Medication errors (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:

Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210506977

Write-up: DEATH; STROKE; EXTRA DOSE ADMINISTERED; This spontaneous report received from a consumer via social media concerned a male of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient administered extra dose and had a bad stroke within 12 hours of the 2nd dose. Later, the patient died. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of death on an unspecified date, and the outcome of extra dose administered was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: V0: 20210506977- Covid-19 vaccine ad26.cov2.s- Death, Stroke. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: STROKE

VAERS ID: 1320104 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021486645

Write-up: dropped dead; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that a patient of unspecified gender in 20s received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection) via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as a single dose for covid-19 immunization. The patient medical history was not reported. The patient''s concomitant medications were not reported. The reporter reported that someone in town in their 20s that had the Pfizer vaccine in the morning and dropped dead in the afternoon for no reason. Between all 3, there have been quite a bit of deaths in the last few months. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: dropped dead

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VAERS ID: 1320235 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-15

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Allergy
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021498273

Write-up: died from Pfizer vaccine of anaphylactic reaction; This is a spontaneous report from a non-contactable consumer. A 60-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as unknown, single for Covid-19 immunisation. Medical history included hypersensitivity (there was something he was allergic to it''s what is used in contrast imaging). Concomitant medications were not reported. The patient had an anaphylactic reaction to the vaccination and through the anaphylactic reaction he passed away. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information about lot/batch number has been requested.; Reported Cause(s) of Death: Anaphylactic reaction

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VAERS ID: 1320290 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-15

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Dyspnoea, Pulmonary oedema
SMQs:, Cardiac failure (narrow), Anaphylactic reaction (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COPD
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC2021504517

Write-up: Lungs filled up with fluid; could not breath after the second dose; This a spontaneous report from a contactable consumer via a Pfizer sales representative. An 89-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as 2nd dose, single for COVID-19 immunization. Medical history included chronic obstructive pulmonary disease (COPD). The patient''s concomitant medications were not reported. On an unspecified date, the patient died due to lungs filled with fluid and could not breath after the second dose. It was not reported if an autopsy was performed. Information on the batch/lot number has been requested.; Reported Cause(s) of Death: Lung filled up with fluid; could not breath after the second dose

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VAERS ID: 1320319 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Product use issue
SMQs:, Medication errors (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021507396

Write-up: Girl''s family who got vaccinated died along with their dog due to have been being exposed to the members who got vaccinated; 11 year old, pre-period, girls family who got

vaccinated; This is a spontaneous report from a contactable consumer. A 11-years-old patient of an unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The reporter reported that 11 year old, pre-period, girls family who got vaccinated died along with their dog due to have been being exposed to the members who got vaccinated. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot/batch number has been requested; Reported Cause(s) of Death: Girl"s family who got vaccinated died along with their dog due to have been being exposed to the members who got vaccinated

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VAERS ID: 1322554 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210516695

Write-up: DEATH; This spontaneous report received from a company representative via social media concerned a female patient of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, Expiry: Unknown) dose not reported, 1 total, on an unspecified date, for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case.

No concomitant medications were reported. In 2021, after 2 weeks of vaccination, the patient was deceased (unknown cause). It was unknown if the autopsy was performed or not. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The putcome of the event was fatal. This report was serious (Death). This case, from the same reporter is linked to 20210517056.; Sender"s Comments: V0; 20210516695-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1322556 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident, Death
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210516928

Write-up: DEATH; STROKE; This spontaneous report received from a consumer concerned eight patients with unknown race and ethnicity. The patients" weight, height, and medical history were not reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has

been requested. No concomitant medications were reported. On an unspecified date, the patients had stroke and died after getting the Janssen Covid-19 vaccine, the patients died from unknown cause of death and it was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0.20210516928-covid-19 vaccine ad26.cov2.s-Death,Stroke. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1322563 (history)
Form: Version 2.0
Age:
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210522533

Write-up: BLOOD CLOTS; This spontaneous report received from a consumer who reported hearing a news report concerned a male of unspecified age. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number not reported, expiry not reported frequency one total, dose, therapy start date were not reported administered for prophylactic vaccination. The batch number was not reported and has been requested.No

concomitant medications were reported. On an unspecified date, the patient had blood clots. On an unspecified date in 2021, the patient was died due to blood clots. it was not reported whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died due to blood clot on an unspecified date in 2021. This report was serious (Death). This case, from the same reporter is linked to 20210523500.; Sender"s Comments: V0. 20210522533-COVID-19 VACCINE AD26.COV2.S-Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOTS

VAERS ID: 1326271 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COPD
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021538103

Write-up: Cardiac arrest; This is a spontaneous report from a contactable consumer reporting for her uncle. A male patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date in 2021, at single dose, for COVID-19 immunization. Medical history included COPD. Concomitant medications were not reported. A week after the patient got his COVID shot, he was dead, he had a cardiac arrest on an unspecified date in 2021, with fatal outcome. This just happened last month. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested. ; Reported Cause(s) of Death: Cardiac arrest

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VAERS ID: 1327525 (history)
Form: Version 2.0
Age: 82.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-02-06
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9264 / 1	LA / SYR

Administered by: Other **Purchased by:** ?
Symptoms: Death, Fatigue, Loss of consciousness, Pain in extremity, Seizure, Unresponsive to stimuli
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), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (narrow), Tendinopathies and ligament disorders (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-11
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: unknown
Current Illness: none
Preexisting Conditions: unknown
Allergies: none known

Diagnostic Lab Data:
CDC Split Type:

Write-up: She received her first dose on 2/6/21 around 11:30 a.m. On 2/7/21 she was a bit tired and her arm was a little sore. Sometime after the afternoon of 2/7/21 and the morning of 2/8/21 she was found unconscious and unresponsive on the floor. As her dinner from the evening of 2/7/21 was still on the table unfinished and she was fully dressed we assume that whatever caused her to end up on the floor unconscious happened on 2/7/21 She was taken to the emergency room and it was determined that she was not going to survive. She never regained consciousness and was having seizures all the way to the hospital in the back of the ambulance. She was made comfortable and passed on 2/11/21.

VAERS ID: 1329078 (history)
Form: Version 2.0
Age: 45.0
Sex: Female
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-19

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	- / UNK	- / -
UNK: VACCINE NOT SPECIFIED (NO BRAND NAME) / UNKNOWN MANUFACTURER	MM248M0101 / UNK	- / OT

Administered by: Other **Purchased by:** ?
Symptoms: Vaccination complication
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: OPSUMIT; WARFARIN
Current Illness: Primary pulmonary hypertension
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210521894

Write-up: REACTION TO COVID VACCINE; This spontaneous report received from a patient concerned a 45 year old female of unspecified race and ethnicity. The patient''s weight was 88.2 kilograms, and height was not reported. The patient''s concurrent conditions included primary pulmonary hypertension. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number: unknown, expiry: unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. The patient received epoprostenol sodium (form of admin unknown, intravenous, batch number: MM248M0101 expiry: 30-JUN-2022) 1.5 mg, frequency, and therapy dates were not reported for drug used for unknown indication. Concomitant medications included macitentan and warfarin for drug used for unknown indication. On an unspecified date, the patient experienced reaction to covid vaccine, and was hospitalized on 07-MAY-2021. On an unspecified date, the patient died from reaction to covid vaccine. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable; and dose of epoprostenol sodium was not changed. This report was serious (Death, and Hospitalization Caused / Prolonged).; Sender''s Comments: V0. 20210521894-COVID-19 VACCINE AD26.COV2.S-REACTION TO COVID VACCINE. This event(s) is considered not related. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: REACTION TO COVID VACCINE

VAERS ID: 1329662 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210530454

Write-up: BLOOD CLOTS; This spontaneous report received from a consumer concerned 3 women with unknown race and ethnicity . The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry was unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date,3 women died from blood clots after getting (Janssen) covid 19 vaccine. It was unknown whether autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0 : 20210530454-COVID-19 VACCINE AD26.COV2.S-Blood clots . This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1332466 (history)
Form: Version 2.0
Age:
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-20

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

Administered by: Other **Purchased by:** ?
Symptoms: Pulmonary embolism
SMQs:, Embolic and thrombotic events, venous (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Routine health maintenance (Very healthy, athletic and a marathon runner.); Comments: marathon runner, no known medical history; very healthy Mother dies last year of aneurysm. Patient''s dad is in his 80''s is

very healthy.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210533144

Write-up: PULMONARY EMBOLISM; This spontaneous report received from a consumer via a company representative concerned a 63 year old male, race and ethnicity unspecified. The patient''s height, and weight were not reported. The patient''s medical history was not reported. The patient was very healthy, athletic and a marathon runner. The patient''s mother died last year of an aneurysm. The patient''s dad is in his 80''s is very healthy. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the subject developed pulmonary embolism causing death six days after receiving the vaccination. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died on an unspecified date. This report was serious (Death).; Sender''s Comments: V0: 20210533144-covid-19 vaccine ad26.cov2.s- This case concerns to a 63 year old male. Pulmonary embolism. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: PULMONARY EMBOLISM

VAERS ID: 1332839 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Illinois
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-20

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Drug ineffective, SARS-CoV-2 test, SARS-CoV-2 test positive
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Chronic lymphocytic leukemia (Since 2018 or 2019)
Allergies:
Diagnostic Lab Data: Test Name: covid test; Test Result: Positive
CDC Split Type: USPFIZER INC2021541162

Write-up: died; tested positive for covid; tested positive for covid; This is a spontaneous report from a non-contactable consumer. This male patient of unspecified age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunisation. Medical history included chronic lymphocytic leukemia untreated since 2018 or 2019. Concomitant medications were not provided. It was reported that on unspecified date, patient got BNT162B2 and later tested positive for covid and died. It was not reported if an autopsy was performed. Lab data included covid test was positive on unspecified date. The outcome of the event died was fatal, while of the other events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: died

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VAERS ID: 1332850 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 2021-02-01
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-20

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Pulmonary embolism, SARS-CoV-1 test
SMQs:, Embolic and thrombotic events, venous (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19 (resolved one month before receiving vaccine)
Allergies:
Diagnostic Lab Data: Test Name: Covid_19 Test; Test Result: Positive ; Comments: before vaccine; Test Date: 202101; Test Name: Covid_19 Test; Test Result: Negative ; Comments: one month before receiving vaccine
CDC Split Type: USPFIZER INC2021547314

Write-up: Pulmonary embolism; This is a spontaneous report received from a Pfizer sponsored program, received by a contactable consumer (patient"s relative). A 91-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Lot number not provided), via an unknown route, in Feb2021 at single dose for COVID-19 immunization, administered at facility. Relevant medical history included COVID positive from an unknown date and resolved in Jan2021 (COVID test negative one moth before receiving vaccine). No relevant concomitant medications were provided. Patient"s relative stated that she died 10 days after receiving the COVID vaccine. The cause of death was not known but caller stated it could be due to pulmonary embolism. The facility did not wait for 3 months before giving the vaccine. Caller stated it should be prominent that patient should wait for 3 months after being tested positive for COVID and receiving the vaccine. It was unknown if an autopsy was performed. Information about Lot/Batch number has been requested.; Reported Cause(s) of Death: pulmonary embolism

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VAERS ID: 1332962 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Montana
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-20

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Platelet count, Platelet count decreased
SMQs:, Haematopoietic thrombocytopenia (narrow), Systemic lupus erythematosus (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Routine health maintenance; Comments: Does the patient have any allergies? : Not asked Was there drug abuse or illicit drug usage? :Unknown Does the patient consume alcohol? :Unknown Does the patient smoke? : Unknown
Allergies:
Diagnostic Lab Data: Test Name: Platelet count; Result Unstructured Data: low (no values provided)
CDC Split Type: USJNJFOC20210526587

Write-up: DEATH; LOW PLATELET; This spontaneous report received from a consumer who reported she had seen a post approximately one month ago concerning a male of unspecified age. The patient''s height, and weight were not reported. The patient''s concurrent conditions included good health. The patient''s medical history was not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: Unknown) dose and start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the subject experienced low platelet and died on an unspecified date. Also conflictingly reported the cause of death was unknown (adverse events captured as Low Platelet and Death). It was not reported if an autopsy was performed. As per the reporter, the patient was an emergency (ER) physician who died after receiving the vaccine due to low platelets. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died on an unspecified date. This report was serious (Death).; Sender''s Comments: V0: 20210526587-covid-19 vaccine ad26.cov2.s -Death, Low platelets. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: LOW PLATELET

VAERS ID: 1340228 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Washington
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-22

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210536600

Write-up: DEATH; This spontaneous report received from a consumer concerned four female patients. The patient''s weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total dose was administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company was unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On unspecified date in 2021, the patients died from unknown cause of death after vaccination. The autopsy details were not provided. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210320422.; Sender''s Comments: V0- 20210536600-Covid-19 vaccine ad26.cov2.s-death.This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1345714 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-25

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210538222

Write-up: HEART ATTACK; This spontaneous report received from a consumer via a company representative, concerned a female patient of unspecified age, race and ethnic origin. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration was not reported, batch number: unknown and expiry: unknown) dose and therapy start date were not reported, 1 total, administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, in 2021 (11 hours post vaccination), the patient died due to heart attack. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of heart attack was fatal. This report was serious (Death).; Sender''s Comments: V0 20210538222-covid-19 vaccine ad26.cov2.s-Myocardial infarction. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: HEART ATTACK

VAERS ID: 1345716 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-25

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210538476

Write-up: DEATH; This spontaneous report received from a consumer via social media concerned a patient of unspecified age and sex. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, frequency one total ,start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the reporter reported that one patient died post vaccination from unknown cause of death. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died due to unknown cause on an unspecified date. This report was serious (Death). This case, from the same reporter is linked to 20210536753.; Sender''s Comments: V0: 20210538476-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1345770 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	UNKNOWN / UNK	- / -
COVID19: COVID19 (COVID19 (UNKNOWN)) / UNKNOWN MANUFACTURER	- / UNK	- / -
UNK: VACCINE NOT SPECIFIED (NO BRAND NAME) / UNKNOWN MANUFACTURER	- / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210541733

Write-up: DEATH; This spontaneous report received from a consumer by a other manufacturer company (Pfizer Inc.) via social media post, was received on 14-MAY-2021 and concerned multiple patients (more than 3000). The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown) dose, start therapy date were not reported, frequency 1 total dose administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. Non-company suspect vaccine included: MRNA 1273 (form of admin, route of admin, and batch number were not reported), dose, start therapy date were not reported for an unspecified indication; and BNT 162 (form of admin, route of admin, and batch number were not reported), dose, start therapy date were not reported for an unspecified indication. No concomitant medications were reported. It was reported that on an unspecified date, more than 3000 patients died from vaccine. The cause of death was unknown. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210541733-covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1345808 (history)
Form: Version 2.0
Age: 67.0
Sex: Female
Location: New York
Vaccinated: 2021-02-28
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	001121A / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Abdominal pain, Acute kidney injury, Anaemia, Asthenia, Azotaemia, Blood culture, Blood gases, Blood pressure measurement, Blood test, Body temperature, Brain

natriuretic peptide, Chest X-ray, Chest pain, Computerised tomogram, Dyspnoea, Echocardiogram, Heart rate, Hyperkalaemia, Hyperphosphataemia, Hyperuricaemia, Hyponatraemia, Lethargy, Malnutrition, Metabolic acidosis, Metabolic encephalopathy, Multiple organ dysfunction syndrome, Nausea, Oxygen saturation, Prothrombin time, Respiratory rate, SARS-CoV-2 test, Troponin, Urine analysis, Waist circumference increased
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Anaphylactic reaction (broad), Acute pancreatitis (broad), Haematopoietic erythropenia (broad), Lactic acidosis (broad), Hyperglycaemia/new onset diabetes mellitus (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), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (narrow), Noninfectious meningitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow), Hyponatraemia/SIADH (narrow), Cardiomyopathy (broad), Lipodystrophy (broad), Chronic kidney disease (narrow), Tumour lysis syndrome (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypoglycaemia (broad), Dehydration (broad), Sepsis (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-03-13
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: Acute renal failure (Due to Foley catheter placement); Breast cancer stage IV; COPD; GERD; Hypertension; Metastatic bone disease prophylaxis (Lumbar spine pelvis and proximal femurs); Peritoneal carcinomatosis; Tobacco user; Transaminases increased
Preexisting Conditions: Medical History/Concurrent Conditions: Pneumonitis
Allergies:
Diagnostic Lab Data: Test Date: 20210312; Test Name: Blood culture; Test Result: Negative ; Result Unstructured Data: Negative; Test Date: 20210312; Test Name: VBG; Result Unstructured Data: Compensated mixed respiratory and Metabolic alkalosis; Test Date: 20210312; Test Name: Blood pressure measurement; Test Result: Inconclusive ; Result Unstructured Data: 96/52 mmHg; Test Date: 20210312; Test Name: Blood pressure measurement; Test Result: Inconclusive ; Result Unstructured Data: 100/50 mmHg; Test Date: 20210312; Test Name: Blood pressure measurement; Test Result: Inconclusive ; Result Unstructured Data: 88/58 mmHg; Test Date: 20210312; Test Name: Blood work; Result Unstructured Data: WBC-2.9 Normal, Hg-Normal, Platelet count-Normal; Test Date: 20210312; Test Name: Body temperature; Test Result: Inconclusive ; Result Unstructured Data: 36.1degrees Celsius; Test Date: 20210312; Test Name: Body temperature; Test Result: Inconclusive ; Result Unstructured Data: 35.9 degrees Celsius; Test Date: 20210312; Test Name: Body temperature; Test Result: Inconclusive ; Result Unstructured Data: 36.2degrees Celsius; Test Date: 20210312; Test Name: BNP; Result Unstructured Data: normal; Test Date: 20210312; Test Name: Chest X-ray; Result Unstructured Data: No acute thoracic pathology; Test Date: 20210312; Test Name: CT of chest/abdomen/pelvis; Result

Unstructured Data: revealed a right sided pleural effusion bony metastatic disease and hepatic metastatic disease, as well as small amount of ascites in the upper abdomen; Test Date: 20210312; Test Name: Echocardiogram; Test Result: Inconclusive ; Result Unstructured Data: Ejection fraction 60-65% and otherwise no significant findings; Test Date: 20210312; Test Name: Heart Rate; Test Result: Inconclusive ; Result Unstructured Data: 92; Test Date: 20210312; Test Name: Heart Rate; Test Result: Inconclusive ; Result Unstructured Data: 86; Test Date: 20210312; Test Name: Heart Rate; Test Result: Inconclusive ; Result Unstructured Data: 90; Test Date: 20210312; Test Name: SpO2; Test Result: Inconclusive ; Result Unstructured Data: 96 percent; Test Date: 20210312; Test Name: SpO2; Test Result: Inconclusive ; Result Unstructured Data: 93 percent; Test Date: 20210312; Test Name: SpO2; Test Result: Inconclusive ; Result Unstructured Data: 98 percent; Test Date: 20210312; Test Name: Prothrombin time; Result Unstructured Data: normal; Test Date: 20210312; Test Name: Respiratory rate; Test Result: Inconclusive ; Result Unstructured Data: breaths per minute; Test Date: 20210312; Test Name: Respiratory rate; Test Result: Inconclusive ; Result Unstructured Data: breaths per minute; Test Date: 20210312; Test Name: Respiratory rate; Test Result: Inconclusive ; Result Unstructured Data: breaths per minute; Test Date: 20210312; Test Name: Covid-19 test; Test Result: Negative ; Result Unstructured Data: Negative; Test Date: 20210312; Test Name: Troponin; Result Unstructured Data: normal; Test Date: 20210312; Test Name: Urinalysis; Result Unstructured Data: normal
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Shortness of breath increasing; lethargy; Hyperkalemia; Acute kidney injury; metabolic encephalopathy; Uremia; metabolic acidosis; protein calorie malnutrition; Hyperuricemia; Hyperphosphatemia; Hyponatremia; Anemia; Substernal chest pain aggravated; increasing abdominal girth; Abdominal Pain; Weakness; Nausea; multiorgan failure; This spontaneous case was reported by a physician assistant (subsequently medically confirmed) and describes the occurrence of MULTIPLE ORGAN DYSFUNCTION SYNDROME (multiorgan failure), DYSPNOEA (Shortness of breath increasing), LETHARGY (lethargy), HYPERKALAEMIA (Hyperkalemia), ACUTE KIDNEY INJURY (Acute kidney injury), METABOLIC ENCEPHALOPATHY (metabolic encephalopathy) and AZOTAEMIA (Uremia) in a 67-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 001121A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient"s past medical history included Breast cancer stage IV since March 2019, COPD since an unknown date, GERD since an unknown date, Hypertension since an unknown date, Tobacco user since an unknown date, Transaminases increased since an unknown date, Peritoneal carcinomatosis since an unknown date and Metastatic bone disease prophylaxis (Lumbar spine pelvis and proximal femurs) since an unknown date. Concurrent medical conditions included Acute renal failure (Due to Foley catheter placement) and Pneumonitis. On 28-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced MULTIPLE ORGAN DYSFUNCTION SYNDROME (multiorgan failure) (seriousness criteria death and medically significant), DYSPNOEA (Shortness of breath increasing) (seriousness criterion hospitalization), LETHARGY (lethargy) (seriousness criterion hospitalization), HYPERKALAEMIA (Hyperkalemia) (seriousness criterion medically significant), ACUTE KIDNEY INJURY (Acute kidney injury) (seriousness criterion medically significant), METABOLIC ENCEPHALOPATHY (metabolic encephalopathy) (seriousness criterion medically significant), AZOTAEMIA (Uremia) (seriousness criterion medically significant), METABOLIC ACIDOSIS (metabolic acidosis), MALNUTRITION (protein calorie malnutrition), HYPERURICAEMIA (Hyperuricemia), HYPERPHOSPHATAEMIA (Hyperphosphatemia), HYPONATRAEMIA (Hyponatremia), ANAEMIA (Anemia), CHEST PAIN (Substernal chest pain aggravated), WAIST CIRCUMFERENCE INCREASED (increasing abdominal girth), ABDOMINAL PAIN (Abdominal Pain), ASTHENIA (Weakness) and NAUSEA (Nausea). The patient was hospitalized on 09-Mar-2021 due to DYSPNOEA and LETHARGY. The patient died on 13-Mar-2021. The reported cause of death was

Multiorgan failure. It is unknown if an autopsy was performed. At the time of death, DYSPNOEA (Shortness of breath increasing), LETHARGY (lethargy), HYPERKALAEMIA (Hyperkalemia), ACUTE KIDNEY INJURY (Acute kidney injury), METABOLIC ENCEPHALOPATHY (metabolic encephalopathy), AZOTAEMIA (Uremia), METABOLIC ACIDOSIS (metabolic acidosis), MALNUTRITION (protein calorie malnutrition), HYPERURICAEMIA (Hyperuricemia), HYPERPHOSPHATAEMIA (Hyperphosphatemia), HYPONATRAEMIA (Hyponatremia), ANAEMIA (Anemia), CHEST PAIN (Substernal chest pain aggravated), WAIST CIRCUMFERENCE INCREASED (increasing abdominal girth), ABDOMINAL PAIN (Abdominal Pain), ASTHENIA (Weakness) and NAUSEA (Nausea) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 12-Mar-2021, Blood culture: negative (Negative) Negative. On 12-Mar-2021, Blood gases: abnormal (abnormal) Compensated mixed respiratory and Metabolic alkalosis. On 12-Mar-2021, Blood pressure measurement: 96/52 (Inconclusive) 96/52 mmHg, 100/50 (Inconclusive) 100/50 mmHg and 88/58 (Inconclusive) 88/58 mmHg. On 12-Mar-2021, Blood test: normal (normal) WBC-2.9 Normal, Hg-Normal, Platelet count-Normal. On 12-Mar-2021, Body temperature: 36.1 (Inconclusive) 36.1degrees Celsius, 35.9 (Inconclusive) 35.9 degrees Celsius and 36.2 (Inconclusive) 36.2degrees Celsius. On 12-Mar-2021, Brain natriuretic peptide: normal (normal) normal. On 12-Mar-2021, Chest X-ray: normal (normal) No acute thoracic pathology. On 12-Mar-2021, Computerised tomogram: abnormal (abnormal) revealed a right sided pleural effusion bony metastatic disease and hepatic metastatic disease, as well as small amount of ascites in the upper abdomen. On 12-Mar-2021, Echocardiogram: inconclusive (Inconclusive) Ejection fraction 60-65% and otherwise no significant findings. On 12-Mar-2021, Heart rate: 92 heart beats per minute (Inconclusive) 92, 86 heart beats per minute (Inconclusive) 86 and 90 heart beats per minute (Inconclusive) 90. On 12-Mar-2021, Oxygen saturation: 96 (Inconclusive) 96 percent, 93 (Inconclusive) 93 percent and 98 (Inconclusive) 98 percent. On 12-Mar-2021, Prothrombin time: normal (normal) normal. On 12-Mar-2021, Respiratory rate: 16 (Inconclusive) breaths per minute, 18 (Inconclusive) breaths per minute and 19 (Inconclusive) breaths per minute. On 12-Mar-2021, SARS-CoV-2 test: negative (Negative) Negative. On 12-Mar-2021, Troponin: normal (normal) normal. On 12-Mar-2021, Urine analysis: normal (normal) normal. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter considered MULTIPLE ORGAN DYSFUNCTION SYNDROME (multiorgan failure), DYSPNOEA (Shortness of breath increasing), LETHARGY (lethargy), HYPERKALAEMIA (Hyperkalemia), METABOLIC ENCEPHALOPATHY (metabolic encephalopathy), AZOTAEMIA (Uremia), METABOLIC ACIDOSIS (metabolic acidosis), MALNUTRITION (protein calorie malnutrition), HYPERURICAEMIA (Hyperuricemia), HYPERPHOSPHATAEMIA (Hyperphosphatemia), HYPONATRAEMIA (Hyponatremia), ANAEMIA (Anemia), CHEST PAIN (Substernal chest pain aggravated), WAIST CIRCUMFERENCE INCREASED (increasing abdominal girth), ABDOMINAL PAIN (Abdominal Pain), ASTHENIA (Weakness) and NAUSEA (Nausea) to be possibly related. No further causality assessment was provided for ACUTE KIDNEY INJURY (Acute kidney injury). Concomitant medications were not provided. Treatment for the events included proton pump inhibitors, oxygen, Tylenol, and comfort care. Company comment: Based on current available information and the temporal association between product use and the start date of the events a causal relationship cannot be excluded.; Sender''s Comments: Based on current available information and the temporal association between product use and the start date of the events a causal relationship cannot be excluded.; Reported Cause(s) of Death: multiorgan failure

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VAERS ID: 1345844 (history)
Form: Version 2.0
Age:
Sex: Female

Location: Colorado
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Patient died 2 hours after getting the vaccine; This spontaneous case was reported by a non-health professional and describes the occurrence of DEATH (Patient died 2 hours after getting the vaccine) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant products were reported. The reporter was worried about the administration of the vaccine and wanted to know if there was a way for someone to help their community. Reporter had contacted everyone in Public health department of the rural hometown. With the amount of adverse reactions with vaccine reporter was concerned that the vaccine was not being stored properly or not been given properly and further reported that one person died and they were not reporting it because they did not trust the government. A patient died two hours after getting the Moderna vaccine. No treatment information provided. This is a case of sudden death in a female patient who died 2 hours after receiving a dose of vaccine. Very limited information regarding this event has been provided at this time.; Sender"s Comments: This is a case of sudden death in a female patient who died 2 hours after receiving a dose of vaccine. Very limited information regarding this event has been provided at this time.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1345947 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-25

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210545044

Write-up: BLOOD CLOTS; This spontaneous report received from a consumer by a other manufacturer company (Pfizer Inc.) received on14-MAY-2021 and concerned multiple (few) patients. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) frequency 1 total, dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. Reporter stated that Johnson and Johnson vaccine caused blood clots and few people died. It was not reported whether autopsy was performed. On an unspecified date, the patients experienced blood clots. On an unspecified date, the patients died from blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Other Medically Important Condition).; Sender''s Comments: V0; 20210545044-covid-19 vaccine ad26.cov2. s Blood clots. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: BLOOD CLOTS

VAERS ID: 1349042 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-26

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

Administered by: Other **Purchased by:** ?
Symptoms: Brain injury, Cardiac disorder, Cerebrovascular accident, Death, Myocardial infarction, Thrombosis
SMQs:, Myocardial infarction (narrow), Ischaemic central nervous system vascular conditions (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210544901

Write-up: DROPPING DEAD; HEART ATTACKS; BLOOD CLOTS; STROKES; BRAIN DAMAGE; HEART CONDITIONS; This spontaneous report received from a consumer who reported reading from many personal social media accounts which concerned multiple patients of unspecified age and sex. No past medical history or concurrent conditions were reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. It was reported that, the patients were suffering from blood clots, dropping dead, had strokes, heart attacks, heart conditions and brain damage after vaccination. It was also reported that, the patients were perfectly healthy before and now they would never be the same. On an unspecified date, the patient died from unknown cause of death. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patients died of unknown cause on an unspecified date, and the outcome of blood clots, strokes, heart attacks, heart conditions and brain damage was not reported. This report was serious (Death,

and Other Medically Important Condition). This case, from the same reporter is linked to 20210534943.; Sender"s Comments: V0: 20210544901-covid-19 vaccine ad26.cov2.s - Dropping dead, brain damage, blood clots, heart attacks and strokes. This event(s) are considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1349104 (history)
Form: Version 2.0
Age: 30.0
Sex: Female
Location: Nevada
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-26

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Heart rate irregular
SMQs:, Cardiac arrhythmia terms, nonspecific (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Died; irregular heartbeat; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Died) and HEART RATE IRREGULAR (irregular heartbeat) in a 30-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced DEATH (Died) (seriousness criteria death and medically significant) and HEART RATE IRREGULAR (irregular heartbeat) (seriousness criterion medically significant). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy

was performed. At the time of death, HEART RATE IRREGULAR (irregular heartbeat) outcome was unknown. Concomitant medication and treatment information were not reported. Action taken with mRNA-1273 in response to the event was Not Applicable. Company comment:Limited information regarding the events has been provided at this time and a causal relationship cannot be excluded. Cause of death not reported.; Sender"s Comments: Limited information regarding the events has been provided at this time and a causal relationship cannot be excluded. Cause of death not reported.; Reported Cause(s) of Death: Died

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VAERS ID: 1349117 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-26

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	UNKNOWN / 1	- / OT

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Dropped dead of a heart attack; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of MYOCARDIAL INFARCTION (Dropped dead of a heart attack) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an

unknown date, the patient experienced MYOCARDIAL INFARCTION (Dropped dead of a heart attack) (seriousness criteria death and medically significant). The reported cause of death was Heart attack. It is unknown if an autopsy was performed. No concomitant medication was reported. No treatment information was reported. It was reported that patient''s funeral was on 11 May 2021. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. This case was linked to MOD21-090166, US-MODERNATX, INC.-MOD-2021-128463, US-MODERNATX, INC.-MOD-2021-128999 (E2B Linked Report).; Sender''s Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. MOD21-090166: US-MODERNATX, INC.-MOD-2021-128463:Same reporter US-MODERNATX, INC.-MOD-2021-128999:; Reported Cause(s) of Death: Heart attack

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VAERS ID: 1349751 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-26

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210546018

Write-up: DIED; This spontaneous report received from a consumer via a company representative concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not

reported, batch number: unknown, expiry: unknown) dose, start therapy date were not reported, 1 total dose administered for prophylactic vaccination. The batch no was not reported ,The company is unable to performed follow up to request batch /Lot numbers .No concomitant medications were reported. It was reported that on an unspecified date, the patient died from covid-19 vaccine ad26.cov2.s. The cause of death was unknown. it was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death); Sender''s Comments: V0: 20210546018-covid-19 vaccine ad26.cov2. s Death. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1353088 (history)
Form: Version 2.0
Age:
Sex: Female
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-27

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Died in the car on the way home after receiving the second dose of the Moderna COVID-19; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Died in the car on the way home after receiving the second dose of the Moderna COVID-19) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was

reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant medication information not provided. Treatment information not provided. Action taken with mRNA in response to the event/s was not applicable. A patient called on 13MAY2021 to report adverse effects she experienced after receiving the Moderna COVID-19 vaccine. During the call, she reported a serious adverse effect for "another lady" who "died in the car on the way home" after receiving the second dose of the Moderna COVID-19 vaccine. The caller reported that the individual received the second dose of the vaccine. No additional information was provided. The Serious Adverse Event Reporting form has been completed. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. This case was linked to US-MODERNATX, INC.-MOD-2021-128107, MOD21-090166, US-MODERNATX, INC.-MOD-2021-128999 (E2B Linked Report).; Sender"s Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. US-MODERNATX, INC.-MOD-2021-128107: MOD21-090166: US-MODERNATX, INC.-MOD-2021-128999:; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1357434 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-05-28

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210550011

Write-up: DEATHS; BLOOD CLOTS; This spontaneous report received from a consumer who reported reading and seeing on the news concerned a patient of unspecified age, race, ethnic origin and sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number not reported, expiry not reported) frequency one total, dose, therapy start date were not reported administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the consumer stated that he read and saw on the news that this vaccine causes patient''s deaths, and blood clots. On an unspecified date, the patient died from unknown cause of death. It was not reported whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of deaths on an unspecified date, and the outcome of blood clots was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender''s Comments: V0: 20210550011-covid-19 vaccine ad26.cov2.s-deaths, blood clots. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1364314 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021599474

Write-up: Cardiac arrest; This is a spontaneous report received by a contactable consumer. A male patient of unknown age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Lot number not provided), via an unknown route, on unknown date at single dose for COVID-19 immunisation. No relevant medical history and concomitant medications were provided. Recently the patient received the first shot of COVID-19 vaccine and subsequently went into cardiac arrest. He unfortunately passed away one week later. It was unknown if an autopsy was performed. Information about lot/batch number has been requested.; Reported Cause(s) of Death: cardiac arrest

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VAERS ID: 1366828 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-02

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210600235

Write-up: DEATH; This spontaneous report received from a consumer concerned a male of unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died due to unknown cause. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0:20210600235 -COVID-19 VACCINE AD26.COV2.S- Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1366906 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-02

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Passed away; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Passed away) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. The patient''s wife reported that the patient passed away a few weeks after taking the Moderna vaccine. Concomitant medication and treatment information were not reported. Action taken with mRNA-1273 in response to the event was Not Applicable Company comment Very limited information regarding this event/s has been provided at this time. Further information has been requested. Reporter did not allow further contact; Sender''s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1370319 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-03

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Son died 10 days after his first moderna vaccine; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Son died 10 days after his first moderna vaccine) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product was not provided by the reporter. Treatment information was not provided. Action taken with mRNA-1273 in response to the events was not applicable. A male patient of an unknown age who received mRNA-1273 died 10 days after his first dose of vaccine. No medical conditions or conmeds were provided. Very limited information regarding these events has been provided at this time. Further information is being pursued.; Sender"s Comments: A male patient of an unknown age who received mRNA-1273 died 10 days after his first dose of vaccine. No medical conditions or conmeds were provided. Very limited information regarding these events has been provided at this time. Further information is being pursued.; Reported Cause(s) of Death: Son died 10 days after his first moderna vaccine

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VAERS ID: 1373784 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 2021-04-18
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EW161 / 1	LA / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Congenital anomaly, Maternal exposure during pregnancy
SMQs:, Congenital, familial and genetic disorders (narrow), Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021602308

Write-up: Miscarriage on 13May2021/seriousness criterion was reported as "Congenital anomaly/birth defect"; Miscarriage on 13May2021/seriousness criterion was reported as "Congenital anomaly/birth defect"; This is a spontaneous report from a contactable consumer (patient). This is the baby case. A 34-year-old female patient received the 1st single dose of BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, in left arm on 18Apr2021 (Batch/Lot Number: Ew161), and the 2nd single dose, always in the left arm, on 02May2021 (Batch/Lot Number: Ew0171), for COVID-19 immunisation. First and second dose were administered at the age of 34 years old. Relevant medical history was reported as none. The patient"s last menstrual period was on 18Feb2021 and pregnancy due date was 26Dec2021. Concomitant medications included unspecified pre-natal vitamins. Relevant past drug history included sulfa. The patient had not received any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was never diagnosed with COVID-19, and since the vaccination the patient had not been tested for COVID-19. The patient reported having miscarriage on 13May2021, diagnosed after the 2nd shot of BNT162b2. A doctor or other healthcare professional office/clinic visit was needed. The fetus had died sometime between the 1st and 2nd shot based on gestational age. Gestation period when the event was observed was reported as 4 weeks. The patient had to undergo dilation and curettage from which the patient was recovering. The seriousness criterion was reported as "Congenital anomaly/birth defect" (no further information reported about the congenital anomaly). No follow up attempts are needed. No further information expected.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-2021602163 mother/baby case; Reported Cause(s) of Death: Miscarriage; Congenital anomaly; Miscarriage; Congenital anomaly

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VAERS ID: 1373801 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021609860

Write-up: Death; This is a spontaneous report from a non-contactable consumer. An adult male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot Number: Unknown) as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient died on an unspecified date. It was not reported if an autopsy was performed. Follow-up attempts are completed. No further information is expected.; Reported Cause(s) of Death: Death

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VAERS ID: 1373835 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021616779

Write-up: All died within the 5 weeks after there first shot; This is a spontaneous report from a contactable consumer. This consumer reported similar events for six patients. This is the sixth of six reports. An elderly patient of unknown gender received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date of 2021, as single dose for covid-19 immunization. The patient" s medical history and concomitant medications were not reported. Patient was residential home with 5 other elderly residents patient. Reporter stated that they all got vaccinated with the first Pfizer shot (Confirmed as Covid 19 vaccine) and they all died within the 5 weeks after their first shot. By 18Mar2021 these residents in the house would have died. The reporter stated that they should not have died they just died. The reporter doesn" t know if it was a side effect, he knows if it was a reaction but all of the six residents that got vaccinated died within their first vaccination shot. Within the 5 weeks after the first vaccination. Reporter stated they died in their sleep supposedly. They died in their sleep. Outcome of the event was fatal. It was unknown if an autopsy was done. The information on the batch/lot number has been requested.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-2021617030 same reporter/SD/AE, different patients; Reported Cause(s) of Death: died within the 5 weeks after there first shot

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VAERS ID: 1373996 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-04

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident, Death
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210601307

Write-up: DEATH; STROKES; This spontaneous report received from a consumer via social media concerned multiple patients of unspecified gender. The patient"s weight, height, and medical history were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patients died of unknown cause. It was reported that the patients had stroke. Date of death and autopsy details were not reported. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died on an unspecified date and the outcome of strokes was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: V0- 20210601307-covid-19 vaccine ad26.cov2.s- Death, Stroke. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1374022 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: Death; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant medication or treatment were reported by the reporter. Company Comment: This is an invalid case due to no identifiable patient. Also, very limited information regarding the event has been provided at this time. This case was linked to MOD-2021-102410 (Patient Link).; Sender"s Comments: This is an invalid case due to no identifiable patient. Also, very limited information regarding the event has been provided at this time.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1374303 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:

Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Died) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant medications were reported. No treatment information was reported. Action taken with mRNA-1273 (Moderna COVID-19 Vaccine) in response to the event was not applicable. Company comment: Very limited information regarding this patient"s death has been provided at this time.; Sender"s Comments: Very limited information regarding this patient"s death has been provided at this time.; Reported Cause(s) of Death: Death

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VAERS ID: 1380664 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-08

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Dyspnoea, Speech disorder
SMQs:, Anaphylactic reaction (broad), Dementia (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), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:

Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: COVID-19 (admitted to the hospital for COVID); Hospitalization
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210607655

Write-up: DEATH; TROUBLE IN BREATHING; UNABLE TO SPEAK; This spontaneous report received from a patient via a company representative via social media concerned a female of unspecified age. The patient''s height, and weight were not reported. The patient''s medical history included covid-19 infection. The patient was admitted to the hospital with fully recovered Covid. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported, per procedure no follow up will be requested for the case. No concomitant medications were reported. On an unspecified date, immediately after vaccine shot the patient experienced trouble in breathing, and next day patient was unable to speak. On an unspecified date, the subject died from unknown cause of death. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of trouble in breathing and unable to speak was not reported. This report was serious (Death).; Sender''s Comments: V0: 20210607655-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1381213 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021616788

Write-up: All died within the 5 weeks after there first shot; This is spontaneous report from a contactable consumer. This the second case out of six cases. An elderly male patient of unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date of 2021 as single dose for covid-19 immunisation. The patient medical history and the patient''s concomitant medications were not reported. On unknown date the patient died within the 5 weeks after the first vaccination. It was not reported if an autopsy was performed. The patient was in a residential home with 5 other elderly residents. He/she died in sleep. Information related Lot/Batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021617030 same reporter, same vaccine, same adverse event, different patient.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1381214 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021616796

Write-up: all died within the 5 weeks after there first shot; This is spontaneous report from a contactable consumer. This the third case out of six cases. An elderly male patient of unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date of 2021 as single dose for covid-19 immunisation. The patient medical history and the patient''s concomitant medications were not reported. On unknown date the patient died within the 5 weeks after the first vaccination. It was not reported if an autopsy was performed. The patient was in a residential home with 5 other elderly residents. He/she died in sleep. Information related Lot/Batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021617030 same reporter, same vaccine, same adverse event, different patient.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1381215 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:

Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021617301

Write-up: all died within the 5 weeks after there first shot; This is spontaneous report from a contactable consumer. This the fifth case out of six cases. An elderly patient of unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date of 2021 as single dose for covid-19 immunisation. The patient medical history and the patient''s concomitant medications were not reported. On unknown date the patient died within the 5 weeks after the first vaccination. It was not reported if an autopsy was performed. The patient was in a residential home with 5 other elderly residents. He/she died in sleep. Information related Lot/Batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-2021617030 same reporter, same vaccine, same adverse event, different patient.; Reported Cause(s) of Death: all died within the 5 weeks after there first shot

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VAERS ID: 1381218 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021618004

Write-up: All died within the 5 weeks after first shot; This is a spontaneous report from a contactable consumer. This consumer reported similar events for six patients. This is the fourth of six reports. An elderly patient of an unknown gender and age received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration, on an unspecified date of 2021, as single dose for COVID-19 immunization. The patient" s medical history and concomitant medications were not reported. On unknown date, in 2021, the patient died within the 5 weeks after the first vaccination. It was not reported if an autopsy was performed. The patient was in a residential home with 5 other elderly residents. He/she died in sleep. The information on the lot number has been requested.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-2021617030 same reporter/SD/AE, different patients; Reported Cause(s) of Death: All died within the 5 weeks after first shot

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VAERS ID: 1381355 (history)
Form: Version 2.0
Age: 39.0
Sex: Male
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-08

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Patient was healthy.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210611846

Write-up: DEATH; This spontaneous report received from a consumer concerned a 39 year old male. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. Patient was healthy. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) frequency 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. Reporter stated that the patient took the vaccine and then died 3 days later (date unspecified). Reporter did not have any further information to provide. On an unspecified date, the patient died from unknown cause of death. It was unknown whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210611846-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1384711 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Indiana
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-09

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

Administered by: Unknown **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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC2021641564

Write-up: This patient had done the same thing, only he ended up taking pneumonia and dying with it; This is a spontaneous report from a contactable consumer or other non hcp. This is a split case report which was conservatively captured as per the statement "This patient had done the same thing". A 75-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Solution for injection; Lot Number and Expiration date was not reported), via an unspecified route of administration on an unspecified date as unknown, single for covid-19 immunisation. The patient"s medical history and concomitant medications were not reported. The patient experienced pneumonia on an unspecified date. It was reported that, when reporter saw her medical doctor, the day before reporting the event, who was very concerned because she had a recent patient that was about the same age as the reporter, he was 75 years old. The patient had done the same thing, only he ended up taking pneumonia and dying with it. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the event was fatal on an unknown date. Information about Lot/Batch number has been requested.; Reported Cause(s) of Death: This patient had done the same thing, only he ended up taking pneumonia and dying with it

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VAERS ID: 1387787 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-10

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021613680

Write-up: gave my mom the COVID vaccine and now she is dead; This is a spontaneous report from a non-contactable pharmacist. This pharmacist reported on behalf of a pharmacy technician"s mother (the patient). An elderly female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot unknown, unknown if first or second dose) solution for injection intramuscular on an unknown date as a single dose for COVID-19 vaccination. Medical history was not reported. Concomitant medication included unspecified medication but has never taken morphine sulfate. The pharmacy technician stated, " I gave my mom the COVID vaccine and now she was dead." The outcome of the event was fatal. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender"s Comments: The information currently available is limited and does not allow a meaningful case evaluation. However, based on chronological connection to the vaccine a causal relationship between event of death (unknown cause) and BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) cannot be completely excluded. The case will be reevaluated 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.; Reported Cause(s) of Death: gave my mom the COVID vaccine and now she is dead

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VAERS ID: 1391050 (history)
Form: Version 2.0
Age: 96.0
Sex: Female
Location: Massachusetts
Vaccinated: 2021-04-16
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-11

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EW0164 / 1	- / -

Administered by: Public **Purchased by:** ?
Symptoms: Abdominal pain upper, Feeling cold, Headache, Peripheral coldness
SMQs:, Acute pancreatitis (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: ELIQUIS; METOPROLOL; SIMVASTATIN; ISOSORBIDE; ASPIRINE
Current Illness: Blood pressure high (Illness/AE:,High blood pressure Onset Date:Oct2009 Stop Date:Ongoing)
Preexisting Conditions: Medical History/Concurrent Conditions: Cardiac disorder (Illness/AE:Heart issue,High blood pressure Onset Date:Oct2009); Heart attack (Illness/AE:Mild heart attack Pertinent Details:Hospitalised received surgery to insert stent.); Stent placement (Pertinent Details:Hospitalised for high blood pressure;surgery to check heart stent.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021658413

Write-up: feeling cold; stomach pains; coldness was in hands and feet; headache; This is a spontaneous report from a contactable consumer (patient"s daughter). A 96-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, formulation: solution for injection Lot Number: EW0164 and Expiration Date was not reported), via an unspecified route of administration in Arm on 16Apr2021, as 1st dose, single dose for COVID-19 immunization. Medical history included hypertension from Oct2009 and ongoing, cardiac disorder from Oct2009 to an unknown date, myocardial infarction from an unknown date and unknown if ongoing, stent placement from an unknown date and unknown if ongoing. Concomitant medication(s) included apixaban (ELIQUIS); metoprolol (METOPROLOL); simvastatin (SIMVASTATIN); isosorbide (ISOSORBIDE); acetylsalicylic acid (ASPIRINE) all taken for an unspecified indication from an unspecified start date and ongoing. It was reported that within the week prior (Prior to death), she complained of being cold, and had stomach pain, coldness was in hands and feet, she also mentioned headache. The outcome of the events was unknow. Follow up needed, further information has been requested.

VAERS ID: 1394058 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Pennsylvania
Vaccinated: 2021-03-17
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-12

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	037A21B / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

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: USMODERNATX, INC.MOD20212

Write-up: This spontaneous case was reported by a patient family member or friend (subsequently medically confirmed) and describes the occurrence of DEATH (She passed away) and THROMBOSIS (Clots in her leg) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 037A21B) for COVID-19 vaccination. No Medical History information was reported. On 17-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced DEATH (She passed away) (seriousness criteria death, hospitalization and medically significant) and THROMBOSIS (Clots in her leg) (seriousness criteria hospitalization and medically significant). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, THROMBOSIS (Clots in her leg) outcome was unknown. No concomitant medications reported. Description: The patient had clots in her leg and was hospitalized. There was talk of amputation, but she passed away before that occurred. No treatment information provided. Company Comment: Very limited information regarding these events have been provided at this time. Further information cannot be requested.; Sender''s Comments: Very limited information regarding these events have been provided at this time. Further information cannot be requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1398443 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-15

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

Administered by: Other **Purchased by:** ?
Symptoms: Adverse drug reaction
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210619563

Write-up: SEVERE SIDE EFFECTS; This spontaneous report received from a patient concerned a female of unspecified age. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient experienced severe side effects. On an unspecified date, the patient died from severe side effects. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0:20210619563 - Covid-19 Vaccine Ad26.Cov2.S - Severe Side Effects. This event is considered Unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: SEVERE SIDE EFFECTS

VAERS ID: 1398495 (history)
Form: Version 2.0
Age: 74.0
Sex: Male
Location: Illinois
Vaccinated: 2021-01-28
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-15

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	007M20A / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Ageusia, Anosmia, Chills, Myalgia
SMQs:, Rhabdomyolysis/myopathy (broad), Taste and smell disorders (narrow), Eosinophilic pneumonia (broad), Tendinopathies and ligament disorders (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 2021-03-06
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: No medical history information was provided.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Myalgia; Chills; Taste loss; Smell loss; This spontaneous case was reported by a consumer and describes the occurrence of MYALGIA (Myalgia), CHILLS (Chills), AGEUSIA (Taste loss) and ANOSMIA (Smell loss) in a 74-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 030M20A and 007M20A) for COVID-19 vaccination. No medical history information was provided. On 28-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 21-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced MYALGIA (Myalgia) (seriousness criterion death), CHILLS (Chills) (seriousness criterion death), AGEUSIA (Taste loss) (seriousness criterion death) and ANOSMIA (Smell loss) (seriousness criterion death). The patient died on 06-Mar-2021. The cause of death was not reported. It is unknown if an autopsy was performed. No Concomitant medication information was provided. No treatment medication information was provided. Company Comment: This is a case of sudden death in a 74-year-old male patient with unknown medical history died after receiving vaccine. Very limited information has been provided at this time.; Sender''s Comments: This is a case of sudden death in a 74-year-old male patient with unknown medical history died after receiving vaccine. Very limited information has been provided at this time.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1398528 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-15

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Myocardial infarction

SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: heart attack; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of MYOCARDIAL INFARCTION (heart attack) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) unknown. On an unknown date, the patient experienced MYOCARDIAL INFARCTION (heart attack) (seriousness criteria death and medically significant). The reported cause of death was Myocardial infarction. It is unknown if an autopsy was performed. Concomitant medications were not reported. No treatment information was provided. Patient reported a family member died from a heart attack shortly after getting your vaccine. Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Myocardial infarction

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VAERS ID: 1401679 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-16

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210626951

Write-up: DIED; This spontaneous report received from a company representative via Social media concerned a 35-year-old female, unspecified race and ethnicity. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. Post vaccination, after one week, on an unspecified date patient died, cause of death was unknown. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210626951-Covid-19 vaccine ad26.cov2.s-Died. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1401705 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-16

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac disorder
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Cardiac problems after receiving either Pfizer or Moderna vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a nurse and describes the occurrence of CARDIAC DISORDER (Cardiac problems after receiving either Pfizer or Moderna vaccine) in an elderly patient of an unknown gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced CARDIAC DISORDER (Cardiac problems after receiving either Pfizer or Moderna vaccine) (seriousness criterion death). The reported cause of death was Cardiac disorder NOS. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Company Comment: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Sender''s Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Reported Cause(s) of Death: Cardiac disorder NOS

VAERS ID: 1402006 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-16

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210626664

Write-up: DEATH; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry: Not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient died. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died due to unknown cause. This report was serious (Death).; Sender"s Comments: V0; 20210626664- COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1402014 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-16

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210629311

Write-up: DEATH; This spontaneous report received from a consumer via a company representative concerned an adult male. Additional live follow up was received on 14-JUN-2021. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) one total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient was died. The cause of death was not reported. It was unknown whether the autopsy was done or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210629311- covid-19 vaccine ad26.cov2.s-death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1405294 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-17

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac disorder
SMQs:

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Cardiac problems after receiving either Pfizer or Moderna vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a health care professional and describes the occurrence of CARDIAC DISORDER (Cardiac problems after receiving either Pfizer or Moderna vaccine) in an elderly patient of an unknown gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced CARDIAC DISORDER (Cardiac problems after receiving either Pfizer or Moderna vaccine) (seriousness criteria death and life threatening). The reported cause of death was cardiac disorder. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. No relevant concomitant medications were reported. It was reported that the reporter has seen that patients above 60 years of age who developed Cardiac problems after receiving either Pfizer or Moderna vaccine. Two patients even passed away post cardiac problems after vaccination. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not applicable. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender''s Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Reported Cause(s) of Death: Cardiac disorder

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VAERS ID: 1406251 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis

SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210626581

Write-up: BLOOD CLOT; This spontaneous report received from a physician concerned a 21 year old male of unspecified race and ethnicity. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient had blood clot at some point following vaccination and died due to it. It was not reported, if the autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case is linked to 20210456400 and 20210457370 (same reporter). .; Sender"s Comments: V0: 20210626581-covid-19 vaccine ad26.cov2.s-This case concerns a 21 year old male, Blood Clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1406256 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210628286

Write-up: STROKE; This spontaneous report received from a consumer via a company representative via social media platform concerned a female of unspecified age. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose at a frequency of 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient got stroke and patient died from stroke. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0:20210628286-JANSSEN COVID-19 VACCINE Ad26.COV2.S - stroke with fatal outcome- This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: STROKE

VAERS ID: 1409735 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?

Symptoms: Death, Malaise, Myocardial infarction
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021683311

Write-up: died, received the Pfizer vaccine; massive heart attack; she wasn''t feeling well; This is a spontaneous report from a non-contactable consumer or other non hcp. A 74-years-old female patient received second dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; formulation: Solution for injection, Lot Number: UNKNOWN), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient had vaccine on an unspecified date and within one day, the patient stated she wasn''t feeling well. It was reported that she had massive heart attack. The reporter reported that the patient died on an unspecified date and do not know what happened. The patient died on an unspecified date for an unspecified reason. It was unknown weather autopsy was performed or not. The outcome of she wasn''t feeling well and massive heart attack was unknown. Request Name: REQ-347122 Product: PFIZER-BIONTECH COVID-19 VACCINE Question: Has anyone reported Heart attack or stroke? Response: Spoke from attached document: In the all-enrolled population of (total N=43,448), the proportions of participants who reported at least 1 SAE during the time period from Dose 1 to the data cutoff date (November 14, 2020) were 0.6% in the BNT162b2 vaccine group and 0.5% in the placebo group. The most common SAEs in the vaccine group which were numerically higher than in the placebo group were acute myocardial infarction (0.02%), and cerebrovascular accident (0.02%), and in the placebo arm numerically higher than in the vaccine arm were pneumonia (0.03%), atrial fibrillation (0.02%), and syncope (0.02%). Occurrence of SAEs involving system organ classes and specific preferred terms were otherwise balanced between treatment groups, including no imbalance overall in cardiovascular serious adverse events. Offered to email for review, caller declined. No follow-up attempts are possible, No further information is expected.

VAERS ID:	1410930 (history)
Form:	Version 2.0
Age:	83.0
Sex:	Male
Location:	Kentucky

Vaccinated: 2021-06-10
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	004C21A / 2	RA / IM

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-06-14
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Eliquis
Current Illness:
Preexisting Conditions:
Allergies: NKA
Diagnostic Lab Data: unknown
CDC Split Type:

Write-up: death, June 14, 2021

VAERS ID: 1412223 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210627096

Write-up: BLOOD CLOT; This spontaneous report received from a patient via social media via a company representative concerned a patient of unspecified age and sex. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose, start therapy date were not reported for prophylactic vaccination.The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient had blood clot and died.The cause of death is blood clot. It was unknown whether an autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210627090.; Sender"s Comments: V0: 20210627096-covid-19 vaccine ad26.cov2.s-blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1412230 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-19

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebral haemorrhage
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhagic central nervous system vascular conditions (narrow)

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 2021-06-10
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210628293

Write-up: BRAIN BLEED; This spontaneous report received from a consumer via social media via a company representative concerned a male of unspecified age. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: Unknown) dose, start therapy date were not reported, 1 total, administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced brain bleed. On Thursday 10-JUN-2021, the patient died from brain bleed. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210628379 and 20210628386.; Sender"s Comments: V0-20210628293-Covid-19 vaccine ad26.cov2.s - BRAIN BLEED . This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BRAIN BLEED

VAERS ID: 1413234 (history)
Form: Version 2.0
Age: 96.0
Sex: Female
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-20

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and

venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-05-25
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: TBD
Current Illness: Stomach cancer
Preexisting Conditions: TBD
Allergies: TBD
Diagnostic Lab Data: March 2021
CDC Split Type:

Write-up: Multiple blood clots that led to death on May 25, 2021.

VAERS ID: 1413724 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-21

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

Administered by: Other **Purchased by:** ?
Symptoms: COVID-19, Death
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:

Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210629346

Write-up: PASSED AWAY; COVID-19; This spontaneous report received from a consumer via a company representative via social media concerned a female of unspecified age. Initial information was processed with the additional information received from central complaint vigilance on 15-JUN-2021. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient got COVID-19 and passed away. The patient died from unknown cause of death. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of COVID-19 was not reported. This report was serious (Death). This report was associated with product quality complaint: 90000182715. The suspected product quality complaint has been confirmed to be voided (did not meet PQC criteria) based on the PQC evaluation/investigation performed. This case, from the same reporter is linked to 20210635272.; Sender"s Comments: V0; 20210629346-covid-19 vaccine ad26.cov2.s ? Passed away. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1413736 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Myocarditis
SMQs:, Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210632922

Write-up: SUDDEN CARDIAC DEATH CAUSED BY MYOCARDITIS; This spontaneous report received via social media from a patient via a company representative concerned a 30 year old male. The patient''s weight, height and medical history were not reported. No past medical history or concurrent conditions were reported. On an unspecified date, the patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: not reported) 1 total dose administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, after eight days of vaccination, the patient died of sudden cardiac death caused by myocarditis. The reporter figured all this in the news and thought it could be related. The cause of death was myocarditis. It was unknown whether an autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210632922-covid-19 vaccine ad26.cov2.s-sudden cardiac death caused by myocarditis. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: MYOCARDITIS

VAERS ID: 1413743 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-05-01
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210634485

Write-up: DEATH; This spontaneous report received from a consumer via a company representative via social media concerned a female of unspecified age. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, administered on 02-MAY-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date in MAY-2021, she died from unknown cause of death. It was reported that, patient"s obituary news was on paper on 17-MAY-2021. It was unspecified that if the autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0- 20210634485- Covid-19 vaccine ad26.cov2.s-Death.This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1416488 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-22

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, 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? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Passed away; Had seizures 2 days after receiving his first dose; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Passed away) and SEIZURE (Had seizures 2 days after receiving his first dose) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced DEATH (Passed away) (seriousness criteria death and medically significant) and SEIZURE (Had seizures 2 days after receiving his first dose) (seriousness criterion medically significant). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, SEIZURE (Had seizures 2 days after receiving his first dose) outcome was unknown. No concomitant medication information was provided. No treatment medication information was provided. Reporter stated she knows of someone and is not totally sure which vaccine was taken, but says Moderna. Action taken with respect to mRNA-1273 was not applicable. This is a case of a sudden death of a Male patient who reportedly had seizures 2 days after receiving his vaccine. Reporter is unsure of vaccine that patient received and this can be a confounding factor. Very limited information has been provided regarding these events. Most recent FOLLOW-UP information incorporated above includes: On 18-Jun-2021: Sender"s Comments: This is a case of a sudden death of a Male patient who reportedly had seizures 2 days after receiving his vaccine. Reporter is unsure of vaccine that patient received and this can be a confounding factor. Very limited information has been provided regarding these events.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1416708 (history)
Form: Version 2.0
Age:
Sex: Female
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-22

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

Administered by: Other **Purchased by:** ?
Symptoms: Aortic thrombosis

SMQs:, Embolic and thrombotic events, arterial (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-05-29
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210628291

Write-up: BLOOD CLOT IN AORTIC ARTERY; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unk) dose not reported, 1 total administered on 27-MAR-202 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced a massive blood clot in her aortic artery blocking all the blood flow to her legs then, pieces broke off and went to her brain and heart. On 28-MAY-2021 13:45 in afternoon the patient''s life support was stopped. On 29-MAY-2021 at 2:01, 9 weeks after vaccination the patient died from blood clot in aortic artery. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: This spontaneous report from consumer concerns a female patient of unspecified age/race/ethnicity who developed "massive blood clot in her aortic artery blocking all the blood flow to her legs then, pieces broke off and went to her brain and heart" 62 days after receiving Janssen COVID-19 vaccine. The patient died on 63rd day post vaccination. No other details reported. The information available precludes a complete and meaningful assessment. Considering the temporal relationship, the events are assessed to have an indeterminate relationship with the vaccination.; Reported Cause(s) of Death: BLOOD CLOT IN AORTIC ARTERY

VAERS ID: 1419747 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-23

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210635272

Write-up: PASSED AWAY; SUSPECTED CLINICAL VACCINATION FAILURE; SUSPECTED COVID-19 INFECTION; This spontaneous report received from a consumer via a company representative via social media concerned a patient of unspecified age, sex, unknown race and ethnicity. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported, expiry: unknown) dose was not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, patient had suspected Covid-19, suspected clinical vaccination failure and patient passed away from unknown cause. It was unknown if the autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was passed away on an unspecified date. The outcome of suspected covid-19 infection and suspected clinical vaccination failure was not reported. This report was serious (Death, and Other Medically Important Condition). This case, from the same reporter is linked to 20210629346.; Sender''s Comments: V0: 20210635272-Covid-19 vaccine ad26.cov2.s-Passed away This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210635272-Covid-19 vaccine ad26.cov2.-Suspected Clinical Vaccination Failure . This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1419824 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Died; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Died) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant medications were provided. It was reported that a person died after receiving a COVID-19 vaccine. No further details were provided. Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1419925 (history)
Form: Version 2.0
Age:
Sex: Unknown

Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021702371

Write-up: patient died recently after getting the vaccine; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient died recently after getting the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information about lot/batch number has been requested.; Reported Cause(s) of Death: patient died recently after getting the vaccine

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VAERS ID: 1419934 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Indiana
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021711228

Write-up: with no health conditions died after the vaccine; This is a spontaneous report from a contactable consumer (patient"s friend). A 25-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. A friend"s 25 year old daughter (the patient) with no health conditions died after the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: with no health conditions died after the vaccine

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VAERS ID: 1420118 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-23

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

Administered by: Other **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210644799

Write-up: DEATH; This spontaneous report received from a parent via a company representative via social media concerned a 30 year old male, unknown race and ethnicity. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported, expiry: unknown) dose was not reported, 1 total, administered on APR-2021 for prophylactic vaccination. The batch number was not reported. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient was died. The patient died from unknown cause of death. It was unknown if the autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210644799-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1422827 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-04-20
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Death

VAERS ID: 1423010 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebral thrombosis, Coma, Death
SMQs:, Hyperglycaemia/new onset diabetes mellitus (broad), Neuroleptic malignant syndrome (broad), Ischaemic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypoglycaemia (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210637172

Write-up: DEATH; COMA; BLOOD CLOTS ON BRAIN; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot number. No concomitant medications were reported. On an unspecified date, the subject experienced death, coma, and blood clots on brain. On an unspecified date, the patient died from blood clots on brain, and coma. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0-20210637172-COVID VACCINE AD26.COV2.S-Death, Coma, Blood clots on brain. This events are considered un-assessable. The events have a compatible/suggestive temporal relationship, are unlabeled, and have unknown scientific plausibility. There is no information on any other factors potentially associated with the events.; Reported Cause(s) of Death: BLOOD CLOTS ON BRAIN; COMA

VAERS ID: 1423016 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Maine
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

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

Administered by: Other **Purchased by:** ?
Symptoms: COVID-19, SARS-CoV-2 test positive, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Rheumatoid arthritis (Treated by immune-suppressing drug (Unspecified))
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data: Test Name: SARS-CoV-2 test positive; Result Unstructured Data: Positive
CDC Split Type: USJNJFOC20210644468

Write-up: DIED FROM COVID-19; SUSPECTED CLINICAL VACCINATION FAILURE; This spontaneous report received from a company representative and a physician via social media concerned a 74 year old female. The patient''s height, and weight were not reported. The patient''s concurrent conditions included rheumatoid arthritis and patient was taking unspecified immune-suppressing medication. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported) dose, start therapy date were not reported for an unspecified indication. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date the patient experienced Covid-19, and suspected clinical vaccination failure. The patient tested positive for Covid-19 four weeks after receiving the vaccine & later died from Covid-19. Her case was extremely rare. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of Covid-19 on an unspecified date and the outcome of suspected clinical vaccination failure was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender''s Comments: V0: 20210644468-COVID-19 VACCINE AD26.COV2.S-DIED FROM COVID-19. This event is considered Unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event. 20210644468-COVID-19 VACCINE AD26.COV2.S-SUSPECTED CLINICAL VACCINATION FAILURE. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS.; Reported Cause(s) of Death: COVID-19

VAERS ID: 1423057 (history)
Form: Version 2.0
Age:
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

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

Administered by: Unknown **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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Polio
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Problems breathing; This spontaneous case was reported by an other health care professional and describes the occurrence of DYSPNOEA (Problems breathing) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. Concurrent medical conditions included Polio. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced DYSPNOEA (Problems breathing) (seriousness criterion death). The reported cause of death was problems breathing. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. Concomitant medications was not provided. Treatment medications was not provided. Action taken with mRNA-1273 in response to events was not applicable. As a young child, patient was in an iron lung. This is a case of fatal Dyspnoea in a male subject with a hx of polio, who died after receiving the second dose of vaccine. Very limited information has been provided at this time.; Sender"s Comments: This is a case of fatal Dyspnoea in a male subject with a hx of polio, who died after receiving the second dose of vaccine. Very limited information has been provided at this time.; Reported Cause(s) of Death: Problems breathing

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VAERS ID: 1423112 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Maryland
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac disorder, Cardiac failure congestive
SMQs:, Cardiac failure (narrow), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021673157

Write-up: congestive heart failure; Heart problem/heart down; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that a 90-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose 1, single and via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose 2, single for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. On an unspecified date, the patient experienced congestive heart failure and heart problem/heart down. The patient had a heart problem after the Pfizer Covid vaccine. The patient had congestive heart failure and was 90 years old. The hospital said for the patient to get the second shot and it shut her heart down and the patient passed away. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on Lot/Batch number has been requested.; Reported Cause(s) of Death: congestive heart failure; Heart problem/heart down

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VAERS ID: 1423115 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021674301

Write-up: died because of the vaccine; This is a spontaneous report from a Non-contactable consumer (patient"s grandson). A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunization. The patient"s medical history and concomitant medications were not reported. The patient died because of the vaccine on an unspecified date. The cause of death was not reported. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: died because of the vaccine

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VAERS ID: 1423128 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Pennsylvania
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021719708

Write-up: Died 3-5 days after getting the Pfizer COVID vaccine; This is a spontaneous report from a contactable consumer. An elderly female patient received bnt162b2 (Pfizer COVID vaccine), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Consumer stated that her aunt, who was going to be 100 years old, died 3-5 days after getting the Pfizer COVID vaccine. Consumer stated that they were healthy people. She stated that she is a transfusion medical specialist. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on lot/batch number has been requested.; Reported Cause(s) of Death: Died 3-5 days after getting the Pfizer COVID vaccine

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VAERS ID: 1426843 (history)
Form: Version 2.0
Age:
Sex: Female
Location: New Hampshire
Vaccinated: 2021-03-26
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-25

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EL9269 / 1	- / -

Administered by: Senior Living **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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021415137

Write-up: chest pain; had some acute shortness of breath; This is a spontaneous report from a contactable nurse (Registered Nurse) via Medical Information Team. This nurse reported for 7 patients. This report is 7 of 7 patient. A female patient of an unspecified age received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection) via an unspecified route of administration on 26Mar2021 (Lot number: EL9269, Expiry date: 01May2021) as single dose for Covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. On an unspecified date, the patient had some acute shortness of breath and chest pain. Reportedly she was calling from the nursing home stated that she was trying multiple times and she needed to speak to someone to report adverse reactions, she had been reporting to the VAERS system since they started giving vaccines in January, stated: "she had summitted probably 30 reports at that point of all different various things in any of the patients even if they were in hospice if they had a vaccine and proceeded to pass away, she done all the reporting. She had a very abnormal large volume of patients that got vaccinated on 16th of April, with the first doses of Pfizer (PFIZER-BIONTECH COVID-19 VACCINE) one specific lot number and she had 7 adverse events in one group of patients out of 30. And it was way too complicated to get that information quickly, so she spoke to somebody as there was a chance that those could be a significant event and needed to tell somebody what was going on". Caller stated: "because all the reports involving one lot it was more suspicious than even all the other reports that she had ever done. It was just one whole group and now she had 3 deaths. She had 3 deaths and have 2 strokes in this group". Caller stated that she would file reports online she just wanted someone to call her back about the side effects and the lot involved. Stated "she got the whole group, who were due to get their second dose on Friday, two days from now, so obviously she not giving it to any of these people there was a 7 of them out of 33. She had 20 staff that have received it the same day she did not have any side effects in any of the staff but definitely little weary at the moment." Offered to forward provide information to safety. Caller provided lot EL9269, Expiry date 01May2021 (stated that it was weird because it was very close to the expiration date). Caller stated: "All those people were dosed on March 26th. Caller stated that she was going to give just basics (in terms of information to start the process) and that she would file a form online. Caller stated: "That day 31 patients received a vaccine and she had 7 patients worth investigating (caller stated that she had that portion written if there was a way to forward. Explained that there was an option to contact through our website but for adverse reports specifically she would refer them to Pfizer safety explained that she also had a fax, but caller declined she already had that information. Verified that she was reporting adverse events (7 patients, gender: 5 females and 2 males). Caller stated "three patients were send out and subsequently passed away in the hospital, one patient with bradycardia, hypotension and she

passed away in the ER, critical labs, she did not even make it one day, we send her out and she passed away in the ER. She had one male patient who had acute stroke she did not have all the details because he was still hospitalized in ICU. She have one (patient) who we sent to the ER hypoxic, hypotensive, short of breath, she ended up passing away in the hospital, her diagnosis there was hypoxia and acute CHF. She had another male patient who had an acute CVA we send him to the hospital he had acute CVA, he had a right artery occlusion, he passed away. She had another female patient who was sent out with shortness of breath and increased confusion, she wound up in the ER with hypoxia and sepsis and she passed away. She was sure that she did not have hospital records, only know what she was told. And then had two others one that was send to the ER with shortness of breath and elevated D-dimer, she actually returned to us her scans were negative, so she was one of those we are not really 100 percent sure, but she did get send out to the ER. And we have another one (female) chest pain, shortness of breath, she was not sent out her D-dimer and her studies that we have done here were within normal limits but definitely had some acute shortness of breath and chest pain. Done troponin and bunch of cardiac labs there, she did not go out. So those were the seven that she had at the moment that were concerning. Outcome of the events was unknown. No follow up attempts are possible. No further information is expected.

VAERS ID: 1426917 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Kentucky
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021683252

Write-up: pass away recently due to blood clots; This is a spontaneous report from a contactable nurse (parent) and consumer. The nurse reported similar events for three patients. This is the first of three reports. A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. The patient pass away recently due to blood clots on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. It was reported that even though the event was outside of reporting timelines & deemed by the PI to be unrelated to study drug. Information on batch and lot number is requested.; Sender"s Comments: Based on available information, a possible contributory role of the subject product, BNT162B2 vaccine, cannot be excluded for the reported event of thrombosis with fatal outcome due to temporal relationship. However, the reported event may possibly represent intercurrent medical condition in this patient. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics including vascular imaging studies, coagulation panel and autopsy results, 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.,Linked Report(s) : PFIZER INC-2021688670 same reporter, event and suspect drug; different patient; PFIZER INC-2021688671 same reporter, event and suspect drug; different patient; Reported Cause(s) of Death: blood clots

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VAERS ID: 1426921 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Kentucky
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-25

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021688670

Write-up: Patient died due to Blood clots; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 3 patients. This is the 2nd of 3 reports. A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as unknown dose, single for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient pass away recently due to blood clots. The agency also stated that all 3 of these nurses had received the Pfizer vaccine. Document submitted for patient even though the event was outside of reporting timelines and deemed by the PI to be unrelated to study drug. It is not reported if autopsy was performed. information about lot/batch number has been requested.; Sender''s Comments: Based from current drug profile, the event thrombosis is assessed as unrelated to suspected drug BNT162B2 . It is difficult to provide possible cause of the event due to limited patient medical information or background.,Linked Report(s) : PFIZER INC-2021683252 same reporter, same drug, same event, different patient;PFIZER INC-2021688671 same reporter, same drug, same event, different patient; Reported Cause(s) of Death: Blood clots

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VAERS ID: 1429304 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-26

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210648705

Write-up: PEOPLE HAD DIED; This spontaneous report received from a consumer via social media concerned multiple patients of unspecified ages with unknown ethnicity and race. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, Expiry: Unknown) dose not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, reporter stated that people had died from vaccine and they were healthy before the vaccination. Patients were died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the event patient had died was fatal. This report was serious (Death). This case, from the same reporter is linked to 20210648176; Sender"s Comments: V0: 20210648705-covid-19 vaccine ad26.cov2.s-People had died . This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1432774 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Myocardial infarction, Thrombosis
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210648060

Write-up: HEART ATTACK; MULTIPLE BLOOD CLOTS; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported ,1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. The reporter stated that, On an unspecified date her mother had multiple blood clots due to which she had heart attack. Patient was on blood thinners and did not survive. It was unknown if autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died from heart attack and had not recovered from multiple blood clots. This report was serious (Death).; Sender"s Comments: V0: 20210648060-covid-19 vaccine ad26.cov2.s -multiple blood clots , heart attack . This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: HEART ATTACK

VAERS ID: 1432778 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210648251

Write-up: DEATH; This spontaneous report received from a consumer via a company representative from social media concerned a female of unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: UNKNOWN) dose, frequency 1 total, start therapy date were not reported administered for prophylactic vaccination. Batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient died. It was reported by consumer that his niece was died due to vaccine. No further information was provided. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210648251-JANSSEN COVID-19 VACCINE Ad26.COV2.S- Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1432788 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Hospitalisation

SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: USJNJFOC20210650080

Write-up: IN THE HOSPITAL 3 DAYS AFTER JANSSEN VACCINE; DEATH (LOST THE FIGHT 3 WEEKS LATER); This spontaneous report received from a consumer via a company representative concerned a female of unspecified age and unknown race and ethnicity. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, one total administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. it was reported that on an unspecified date, the patient was in the hospital fighting for her life 3 days after taking the vaccine. She lost the fight 3 weeks later (death). It was mentioned that the patient believed the propaganda and lost her life. The patient died due to unknown cause. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0:20210650080-COVID-19 VACCINE AD26.COv2.S-In the Hospital 3 days after Jannsen vaccine, Death(Lost the Fight 3 weeks later). This event(s) is considered un-assessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1432800 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210651615

Write-up: DEATH; This spontaneous report received from a consumer concerned a patient of unspecified age and sex. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: UNKNOWN) frequency 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210651615-COVID-19 VACCINE AD26.COV2.S -Death. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1432866 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-29

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Her son has died; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Her son has died) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No relevant concomitant medications were reported. No treatment information was provided. This is a case of death in a male subject who died after receiving first dose of vaccine Very limited information has been provided at this time.; Sender"s Comments: This is a case of death in a male subject who died after receiving first dose of vaccine Very limited information has been provided at this time.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1432900 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-29

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021765175

Write-up: and have a question about a death being reported as possible link to COVID-19 vaccine; This is a spontaneous report from a contactable consumer. An unspecified age and gender patient received an unspecified dose of BNT162B2 via an unspecified route of administration on an unspecified date for COVID-19 immunization. Medical history and concomitant medication were not reported. On an unknown date, the patient experienced death and it was been reported as possible link to COVID-19 vaccine. The reporter stated that was reported a teen dead several days after gotten a vaccine. Since only Pfizer vaccine is available to teens and reporter wanted to know whether have a statement about this. The CDC was investigated any potential link. The outcome of event was fatal. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: and have a question about a death being reported as possible link to COVID-19 vaccine

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VAERS ID: 1435925 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Product contamination, Pyrexia, Septic shock
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Toxic-septic shock conditions (narrow), Drug reaction with eosinophilia and systemic symptoms

syndrome (broad), Sepsis (narrow), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Fatal fevers; Septic shock; Product contamination; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a non-health professional and describes the occurrence of PYREXIA (Fatal fevers), SEPTIC SHOCK (Septic shock) and PRODUCT CONTAMINATION (Product contamination) in an elderly patient of an unknown gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) at an unspecified dose. On an unknown date, the patient experienced PYREXIA (Fatal fevers) (seriousness criterion death), SEPTIC SHOCK (Septic shock) (seriousness criteria death and medically significant) and PRODUCT CONTAMINATION (Product contamination) (seriousness criterion death). The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product use was not provided. 104 Patients received the vaccine. Treatment information was not provided. Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Sender''s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1435953 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-06-30

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

Administered by: Unknown **Purchased by:** ?

Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Death; This spontaneous case was reported by a non-health professional and describes the occurrence of DEATH (Death) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Sender"s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1439574 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210666298

Write-up: DEATH; This spontaneous report received from a consumer via other company Pfizer concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) one total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient was died. The cause of death was not reported. It was unknown whether the autopsy was done or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: 20210666298-Covid-19 vaccine ad26.cov2.s -Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1440065 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Autopsy, Cardiomegaly, Death, Pericardial effusion
SMQs:, Cardiac failure (broad), Systemic lupus erythematosus (broad), Haemodynamic oedema, effusions and fluid overload (narrow), Cardiomyopathy (broad), Drug reaction with

eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: autopsy; Result Unstructured Data: Test Result:enlarged heart and fluid surrounding the heart; Comments: caused by the Covid vaccination
CDC Split Type: USPFIZER INC2021760279

Write-up: died three days after Covid vaccination; Autopsy showed enlarged heart and fluid surrounding the heart; Autopsy showed enlarged heart and fluid surrounding the heart; This is a spontaneous report from a contactable consumer or other non-health care professional in response to mail sent regarding the confirmation of below mentioned query. A 13-year-old male patient received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, Batch/Lot number and Expiration date not reported) via an unspecified route of administration in an unspecified anatomical location on an unspecified date as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient previously received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, Batch/Lot number and Expiration date not reported) via an unspecified route of administration in an unspecified anatomical location on an unspecified date as single dose for COVID-19 immunisation. On an unspecified date, the patient after receiving his second Covid vaccine from Pfizer died three days later. The patient underwent lab tests and procedures which included autopsy: enlarged heart and fluid surrounding the heart caused by the Covid vaccination. The outcome of the events was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: enlarged heart and fluid surrounding the heart

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VAERS ID: 1442349 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-02

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	UNKNOWN / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210666397

Write-up: BLOOD CLOTS; This spontaneous report received from a consumer via a Business partner (Pfizer Inc.) on 25-JUN-2021 concerned three patients of unspecified age and gender. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown and expiry date: unknown) dose was not reported, 1 total, start therapy date was not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, three patients were developed blood clot after vaccination. The patient died from blood clots on an unspecified date. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0:20210666397-Covid-19 vaccine ad26.cov2.s -Blood clots. This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: BLOOD CLOTS

VAERS ID: 1442361 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-02

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	UNKNOWN / UNK	- / -

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210667006

Write-up: DEATH; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age, race and ethnicity. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died from the vaccine due to unknown cause of death and could never got the second shot. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210667006- JANSSEN COVID-19 VACCINE Ad26.COV2.S- Death. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1445751 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-03

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	UNKNOWN / UNK	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Received the Moderna Shot and within 7-8 weeks after passed away; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a patient family member or friend and describes the occurrence of DEATH (Received the Moderna Shot and within 7-8 weeks after passed away) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. The patient"s past medical history included No adverse event. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant medication was reported by reporter. Patient died on an unknown date approximately 7 to 8 weeks after receiving vaccination.; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: unknown casuse of death

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VAERS ID: 1446010 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-03

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

PFIZER/BIONTECH		

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Headache, Loss of consciousness
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), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Hypotonic-hyporesponsive episode (broad), Generalised convulsive seizures following immunisation (broad), Hypoglycaemia (broad)

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021746877

Write-up: death; lost consciousness; headache; This is a spontaneous report from a non-contactable consumer via a regulatory authority. A 33-year-old patient of an unspecified gender received bnt162b2 (BNT162B2), via an unspecified route of administration on an unspecified date at a single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. On an unknown date, the patient experienced lost consciousness, headache, and death. The event, lost consciousness, was life-threatening. The event, death, was serious for death. The clinical course was as follows: the patient was a pilot who had been vaccinated with bnt162b2. A strong headache occurred immediately after starting the flight. Two hours later, the patient almost lost consciousness and made an emergency landing. (withheld) inoculated and killed three pilots. It''s been a fuss, but it doesn''t stop anymore. The patient did not know when it will fall when the patient flies (as reported). The clinical outcome of the events, lost consciousness and headache, was unknown. The clinical outcome of the event, death, was fatal. The patient died on an unspecified date due to an unknown cause of death. It was not reported if an autopsy was performed. No follow-up attempts are possible. No further information is expected.; Reported Cause(s) of Death: death

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VAERS ID: 1446013 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-03

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombocytopenia
SMQs:, Haematopoietic thrombocytopenia (narrow), Systemic lupus erythematosus (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021749751

Write-up: thrombocytopenia; This is a Literature report. The full publication has been requested. A patient of unspecified age and gender received BNT162B2 (lot number unknown), via an unspecified route of administration, on an unknown date at single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient experienced thrombocytopenia and eventual death on an unknown date. The patient died on an unknown date. The cause of death was thrombocytopenia. It was unknown if an autopsy was performed. The outcome of the event was fatal. The recent report of thrombocytopenia and eventual death of a healthy person who received an mRNA-based COVID-19 vaccine has again raised concerns about whether vaccines trigger ITP. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender''s Comments: Based on a very limited information provided in the report, lacking time to onset, history of pre-existing medical conditions, baseline laboratory data, conmeds and course of events, the Company did not consider the event, fatal thrombocytopenia, as secondary to BNT162B2.; Reported Cause(s) of Death: thrombocytopenia

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VAERS ID: 1449426 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown

Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-06

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210703030

Write-up: DIED; This spontaneous report received from a consumer via a company representative via social media concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) one total, dose not reported, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced died. On an unspecified date, the patient was died from unknown cause of death. It was unknown whether the autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0 20210703030-JANSSEN COVID-19 VACCINE-died. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1456637 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Epistaxis
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow)

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: No medical history was reported.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20211

Write-up: healthy woman dying within 48 h of vaccination; blood coming out of her nose; This spontaneous case was reported by a physician and describes the occurrence of DEATH (healthy woman dying within 48 h of vaccination) and EPISTAXIS (blood coming out of her nose) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced DEATH (healthy woman dying within 48 h of vaccination) (seriousness criteria death, medically significant and life threatening) and EPISTAXIS (blood coming out of her nose) (seriousness criteria death and life threatening). The cause of death was not reported. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. No concomitant medications were provided by the reporter. The subject stated that "there are many deaths related to the Covid vaccine and autopsies have been ordered." The subject also stated that many deaths have occurred within 48 hours of the vaccination. He also reported that a healthy young woman was found deceased in her apartment within 48 hours of vaccination. The deceased was found with blood coming out of her nose. No treatment information was provided by the reporter. Very limited information regarding this events has been provided at this time. Details regarding the exact brand of vaccine received, dosing dates, Medical history, concomitant medication, and date/cause of death are required for further evaluation. Further information has been requested.; Sender''s Comments: Very limited information regarding this events has been provided at this time. Details regarding the exact brand of vaccine received, dosing dates, Medical history, concomitant medication, and date/cause of death are required for further evaluation. Further information has been requested.; Reported Cause(s) of Death: Many deaths related to the COVID vaccine

VAERS ID: 1456646 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 2021-03-03
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EN6203 / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Antibody test, COVID-19, Coombs test, Death, SARS-CoV-2 antibody test negative
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Hodgkin''s disease (and they took his spleen out so he is a little immunocompromised.)
Allergies:
Diagnostic Lab Data: Test Date: 20210407; Test Name: IG; Test Result: Negative ; Comments: AbIG Quantitative; Test Date: 20210524; Test Name: IG; Test Result: Negative ; Comments: IGg; Test Date: 2021; Test Name: IG; Test Result: Negative ; Comments: IGgAb, ABIGm; Test Date: 20210407; Test Name: COB TEST; Test Result: Negative ; Test Date: 20210524; Test Name: COB TEST; Test Result: Negative ; Test Date: 2021; Test Name: COB TEST; Test Result: Negative ; Test Date: 20210407; Test Name: COVID-19 test; Test Result: Negative ; Test Date: 20210524; Test Name: COVID-19 test; Test Result: Negative ; Test Date: 20210407; Test Name: SARS test; Test Result: Negative ; Test Date: 20210524; Test Name: SARS test; Test Result: Negative ; Test Date: 2021; Test Name: SARS test; Test Result: Negative
CDC Split Type: USPFIZER INC2021640296

Write-up: passed away; The initial case was missing the following minimum criteria: adverse event. Upon receipt of follow-up information on 01Jul2021, this case now contains all required information to be considered valid. This is a spontaneous report from a contactable consumer (patient) and patient''s wife. A 69-years-old male patient received first dose of BNT162b2

(PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Lot Number: EN6201), via an unspecified route on 10Feb2021 (at the age of 69-years) as dose 1, single and second dose of BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Lot Number: EN6203), via an unspecified route on 03Mar2021 (at the age of 69-years) as dose 2, single for covid-19 immunisation. The patient medical history included Hodgkin''s disease from 1978 (they took his spleen out, so he was a little immunocompromised and they took out his spleen and appendix after diagnosis) and unknown if ongoing. The patient concomitant medications were not reported. On unspecified date, the patient experienced had 3 antibody tests- one that was a 15minutes finger prick and two others that were all 2-3 weeks apart. Patient stated all 3 tests were negative for antibodies and his doctors don''t know why. His doctors said maybe it did not detect antibodies to the vaccine. In 1978, he had Hodgkin''s disease and they took his spleen out, so he was a little immunocompromised. However, the infectious disease doctor said since he got both doses he should still be protected. Reporter was concerned that he didn''t had great immune system when three negative tests came back, he just want to make sure that everybody thinks he was not okay because he was going out over usual instructions because of them. His primary care physician (PCP) said he talked to another doctor and said if he had the vaccine it should be covered but the test came back negative. Reporter stated that the "last test he had was on 24May2021, it was COVID-19 antibody test, SARS test, COB test 2, AbIG Quantitative. This one was done on maybe 10th (unspecified date in 2021) it was rapid test for SARS, COB-2, IGg antibody. That was done on 07Apr201 and stated if Pfizer want, he can give that also, SARS, COVID, COB-2, IGgAb, AbIGm, they all came back negative". Patient''s wife stated her husband passed away and at the time of the vaccination she and her daughter got adverse reactions she would like to fill the form for both her and her daughter. The patient died on unspecified date in 2021 with unknown cause of death and autopsy results was unknown. The outcome of the event was fatal. Follow up needed, further information has been requested.; Reported Cause(s) of Death: passed away

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VAERS ID: 1458799 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Tennessee
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-09

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebrovascular accident
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Egg allergy; Tuberculosis (Tuberculosis in their kidney.)
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: they believe the cause of death was stroke; This spontaneous case was reported by a consumer and describes the occurrence of CEREBROVASCULAR ACCIDENT (they believe the cause of death was stroke) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. Previously administered products included for Product used for unknown indication: Pneumonia vaccine. Past adverse reactions to the above products included Pneumonia with Pneumonia vaccine. Concurrent medical conditions included Egg allergy and Tuberculosis (Tuberculosis in their kidney.). On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced CEREBROVASCULAR ACCIDENT (they believe the cause of death was stroke) (seriousness criteria death and medically significant). The reported cause of death was Stroke. It is unknown if an autopsy was performed. This spontaneous case was reported by a consumer and describes the occurrence of CEREBROVASCULAR ACCIDENT (they believe the cause of death was stroke) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. Previously administered products included for Product used for unknown indication: Pneumonia vaccine. Past adverse reactions to the above products included Pneumonia with Pneumonia vaccine. Concurrent medical conditions included Egg allergy and Tuberculosis (Tuberculosis in their kidney.). On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced CEREBROVASCULAR ACCIDENT (they believe the cause of death was stroke) (seriousness criteria death and medically significant). The reported cause of death was Stroke. It is unknown if an autopsy was performed. The patient was on blood pressure medications. The reporter reports that two days after a family member died two days after they received their first dose of the Moderna COVID-19 vaccine and they believe the cause of death was a stroke. No laboratory data was provided. No Treatment information was provided. Action taken with mRNA-1273 in response to the event was not applicable. Very limited information regarding this event has been provided at this time. No follow up is possible.; Sender''s Comments: Very limited information regarding this event has been provided at this time. No follow up is possible.; Reported Cause(s) of Death: Stroke

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VAERS ID: 1459256 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-07-09

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021827105

Write-up: already has 1 acquaintance who passed the msm and died; This is a spontaneous report from a contactable physician. This physician reported for an unknown patient. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as dose number unknown, single for COVID-19 immunization. The patient medical history and were not reported. The patient experienced already has 1 acquaintance who passed the msm and died on an unspecified date. Reporter stated, First of all, I want to say that I''m going to file an investigation against everything that''s going on. when I chose Pfizer it was because you said the vaccine was good for patients and cancer, chronic diseases. I lived working paying my bills and taking care of my children until the day I got this vaccine. and then all my exams started to change. I know you don''t want to be responsible for anything, because only money matters. already has 1 acquaintance who passed the msm and died. so what does pfizer do to support it? nothing? I''m like 1 vegetable and what are you going to do for me? Somehow I''m going to sue you because a lawyer also finds fault with you for that. very disappointed and living hell and even depression because of my health. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Sender''s Comments: Based on available information and temporal relationship, a possible contributory role of BNT162B2 vaccine cannot be excluded for the reported event of Death. 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.; Reported Cause(s) of Death: Died

VAERS ID: 1459429 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-09

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebral thrombosis
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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Died with a blood clot in her brain; This spontaneous case was reported by a consumer and describes the occurrence of CEREBRAL THROMBOSIS (Died with a blood clot in her brain) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced CEREBRAL THROMBOSIS (Died with a blood clot in her brain) (seriousness criteria death and medically significant). The reported cause of death was died with a blood clot in her brain. It is unknown if an autopsy was performed. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Concomitant product use was not provided by the reporter. No treatment information was provided. Company Comment: Very limited information on this fatal case. However, based on the current available information and assumed temporal association between the use of the unspecified Moderna Vaccine product and the start date of the event, a causal relationship cannot be excluded. Reporter did not allow further contact; Sender''s Comments: Very limited information on this fatal case. However, based on the current available information and assumed temporal association

between the use of the unspecified Moderna Vaccine product and the start date of the event, a causal relationship cannot be excluded.; Reported Cause(s) of Death: Died with a blood clot in her brain

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VAERS ID: 1463018 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-11

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210713544

Write-up: PERISHED APPROX 9 WEEKS POST INJECTION OF YOUR PRODUCT; This spontaneous report received from a company representative via social media and concerned a male patient of unspecified age, race and ethnic origin. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number not reported, expiry not reported frequency one total, dose, therapy start date were not reported administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient perished approximately 9 weeks (2021) post injection and cause of death was unknown. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210713544-covid-19 vaccine ad26.cov2.s -Perished approx. 9 weeks post injection of your product. This event(s) is

considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1463061 (history)
Form: Version 2.0
Age: 13.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-11

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Product administered to patient of inappropriate age
SMQs:, Medication errors (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Died three days after vaccine; 13 year old boy dies three days after the Moderna vaccine; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Died three days after vaccine) in a 13-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced PRODUCT ADMINISTERED TO PATIENT OF INAPPROPRIATE AGE (13 year old boy dies three days after the Moderna vaccine). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, PRODUCT ADMINISTERED TO PATIENT OF INAPPROPRIATE AGE (13 year old boy dies three days after the Moderna vaccine) had resolved. The action taken with mRNA-1273 (Moderna

COVID-19 Vaccine) (Unknown) was unknown. Concomitant product was not provided by the reporter. Treatment information was unknown. Company comment: This is a case of death in a 13-year-old male subject with unknown medical history, who died one day after receiving the vaccine. Very limited information has been provided at this time. Further information has been requested.; Sender"s Comments: This is a case of death in a 13-year-old male subject with unknown medical history, who died one day after receiving the vaccine. Very limited information has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1463262 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-11

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Drug ineffective, SARS-CoV-2 test, Suspected COVID-19
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: PCR Test; Test Result: Negative ; Test Name: PCR Test; Test Result: Negative
CDC Split Type: USPFIZER INC2021785213

Write-up: suspected covid 19; drug ineffective; This is a spontaneous report from a non-contactable consumer (patient daughter) through a Pfizer sponsored program. A male patient of an unspecified age received bnt162b2 (Pfizer-BioNTech Covid-19 Vaccine), via an unspecified route of administration on an unspecified date (Batch/Lot number was not

reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. As per the reporter, her father (orthopedic surgeon) died of covid-19 vaccine on an unspecified date. The patient underwent lab tests and procedures which included had 2 negative PCR test 14 days later. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; batch/lot number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: suspected covid

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VAERS ID: 1466124 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Washington
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-13

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021807766

Write-up: blood clots; This is a spontaneous report from a contactable physician via a sales representative. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Patient died one week after receiving Pfizer COVID vaccine, autopsy

confirmed blood clots. Event took place after use of product. Information on the lot/batch number has been requested.; Sender"s Comments: A possible contributory role of the suspect drug to the reported event "blood clots" cannot be completely excluded based on temporal association. This case will be re-assessed when 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 identifies as part of this review, as well as any appropriate action in response, will be promptly notifies to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.; Reported Cause(s) of Death: blood clots

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VAERS ID: 1469692 (history)
Form: Version 2.0
Age: 55.0
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-14

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	NOT PROVIDED / UNK	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Myocardial infarction, Rheumatoid arthritis
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Arthritis (narrow), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: HUMIRA
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: massive heart attack; Death; rheumatoid arthritis; This spontaneous case was reported by a consumer and describes the occurrence of MYOCARDIAL INFARCTION (massive heart attack), DEATH (Death) and RHEUMATOID ARTHRITIS (rheumatoid arthritis) in a 55-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine)

(batch no. Not provided) for COVID-19 vaccination. Concomitant products included ADALIMUMAB (HUMIRA) for Rheumatoid arthritis. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced MYOCARDIAL INFARCTION (massive heart attack) (seriousness criteria death and medically significant), DEATH (Death) (seriousness criteria death and medically significant) and RHEUMATOID ARTHRITIS (rheumatoid arthritis) (seriousness criteria death and medically significant). The reported cause of death was massive heart attack. It is unknown if an autopsy was performed. No laboratory was not given. No treatment information was provided by reporter. According to, "The caller stated that she does not know if her brother?s friend received a Moderna, Pfizer or J&J vaccine." This a report of dead one day after the second dose of the product in a 55-year-old patient with concomitant Rheumatoid arthritis. Very limited information regarding the event has been provided for inferring causality. Further information has been request. Most recent FOLLOW-UP information incorporated above includes: On 08-Jul-2021: Follow Up information was received 08-JUL-2021 and added concomitant product and event verbatim; Sender"s Comments: This a report of dead one day after the second dose of the product in a 55-year-old patient with concomitant Rheumatoid arthritis. Very limited information regarding the event has been provided for inferring causality. Further information has been request.; Reported Cause(s) of Death: massive heart attack

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VAERS ID: 1481132 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Vaccine breakthrough infection
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: USJNJFOC20210724474

Write-up: BREAKTHROUGH CASE; This spontaneous report received from a health care professional via social media concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose,1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient was hospitalized at the time of death and was a vaccine breakthrough case. On an unspecified date, the patient died. It was unknown autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome was fatal. This report was serious (Death, and Hospitalization Caused / Prolonged). This report was associated with product quality complaint:90000185655; Sender''s Comments: V0; 20210724474 -covid-19 vaccine ad26.cov2- vaccine breakthrough case. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1481184 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210726824

Write-up: DEATH; This spontaneous report received from a consumer via social media company representative concerned a male of unspecified age, race and ethnic origin. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received Covid-19 vaccine ad26.cov2. s (suspension for injection, route of admin not reported, batch number: Unknown expiry: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death. It was reported that, the patient death occurred after getting the vaccine. It was unknown, if an autopsy was performed. The action taken with Covid-19 vaccine ad26.cov2. s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210726824-Covid-19 vaccine ad26.cov2. s -Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1481259 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210730562

Write-up: PASSED AWAY DUE TO THE JOHNSON AND JOHNSON VACCINE; This spontaneous report received from a patient concerned a 4 decade old male. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown), 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient passed away due to the Johnson and Johnson vaccine and cause of death was unknown. It was unspecified if autopsy performed. Reporter just got off the phone with her cousin it was her best friend who passed away. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210730562-covid-19 vaccine ad26.cov2.s-passed away due to the Johnson and Johnson vaccine. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

VAERS ID: 1481771 (history)
Form: Version 2.0
Age: 84.0
Sex: Male
Location: Connecticut
Vaccinated: 2021-07-07
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Arteriosclerosis, Parkinson's disease
SMQs:, Neuroleptic malignant syndrome (broad), Parkinson-like events (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-07-08
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Drug allergy (Allergy to Sinemet)
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Cause of death to be Parkinson"s Disease secondary to ASHD (atherosclerotic heart disease); Atherosclerotic Cardiovascular Disease; This spontaneous case was reported by a nurse and describes the occurrence of PARKINSON"S DISEASE (Cause of death to be Parkinson"s Disease secondary to ASHD (atherosclerotic heart disease)) and ARTERIOSCLEROSIS (Atherosclerotic Cardiovascular Disease) in an 84-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. Concurrent medical conditions included Drug allergy (Allergy to Sinemet). On 07-Jul-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced PARKINSON"S DISEASE (Cause of death to be Parkinson"s Disease secondary to ASHD (atherosclerotic heart disease)) (seriousness criteria death and medically significant) and ARTERIOSCLEROSIS (Atherosclerotic Cardiovascular Disease) (seriousness criterion death). The patient died on 08-Jul-2021. The reported cause of death was parkinson"s disease and Atherosclerotic cardiovascular disease. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. The patient passed away (08Jul2021 at 4:10 PM). The HCP reported that the death certificate showed cause of death to be Parkinson"s Disease secondary to ASHD (atherosclerotic heart disease). No concomitant medication details was provided. No treatment medication details was provided. Very Limited information regarding the events has been provided at this time . Further information has been requested.; Sender"s Comments: Very Limited information regarding the events has been provided at this time . Further information has been requested.; Reported Cause(s) of Death: Parkinson"s Disease; Atherosclerotic Cardiovascular Disease

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VAERS ID: 1481796 (history)
Form: Version 2.0
Age: 26.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-18

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Toxicity to various agents
SMQs:, Anticholinergic syndrome (broad), Drug abuse and dependence (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Toxic effects of Moderna vaccine; This spontaneous case was reported by a consumer and describes the occurrence of TOXICITY TO VARIOUS AGENTS (Toxic effects of Moderna vaccine) in a 26-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced TOXICITY TO VARIOUS AGENTS (Toxic effects of Moderna vaccine) (seriousness criterion death). The reported cause of death was toxic effects of moderna vaccine. It is unknown if an autopsy was performed. No relevant concomitant medications were reported. No treatment information was provided. Company comment: Very limited information regarding this event has been provided at this time. Critical details including medical history, concomitant medication, date of vaccine administration and the event term, and cause of death is required for further assessment.; Sender''s Comments: Very limited information regarding this event has been provided at this time. Critical details including medical history, concomitant medication, date of vaccine administration and the event term, and cause of death is required for further assessment.; Reported Cause(s) of Death: Toxic effects of Moderna vaccine

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VAERS ID: 1483260 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Headache
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021818626

Write-up: Ended up passing away about seven days after she received the second shot of the vaccine; Extreme headache; This is a spontaneous report from a non-contactable consumer. A female patient (reporter"s mother) of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on an unspecified date as DOSE 2, SINGLE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient previously received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), for covid-19 immunisation. Patient actually took both of the vaccines and she experienced extreme headache after she received her second vaccine and she ended up passing away, she ended up passing away about seven days after she received the second shot of the vaccine. The outcome of headache was unknown, while other event was fatal. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Ended up passing away about seven days after she received the second shot of the vaccine

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VAERS ID: 1483624 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebrovascular accident, Investigation
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)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Organ test; Result Unstructured Data: Test Result:Normal Results; Comments: she was perfectly fine
CDC Split Type: USPFIZER INC2021873485

Write-up: Stroke; This is a spontaneous report received from a contactable consumer. This consumer reported for a female patient (reporters friend"s mom) that an unspecified age elder female patient received second dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot number and expiration date was not reported), via unspecified route of administration on an unspecified date in 2021 as dose 2, single for COVID-19 immunization. The patient medical history and concomitant medication was not reported. The patient previously took first dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot number and expiration date was not reported), via unspecified route on an unspecified date as single dose for COVID-19 immunization. On an unspecified date patient died she had a stroke, and they were trying to say that because she was older, she just passed her organ test, and she was perfectly fine and 2 days later after her second shot she had a stroke (Not clarified hence split not made) and it was like they keep on and understood that there were multiple reasons that someone could have a stroke specially if patient was older. The outcome of evet was fatal Information on the lot/batch number has been requested. Additional information is requested.; Reported Cause(s) of Death: stroke

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VAERS ID: 1483629 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 2021-03-29
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-18

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Atrial fibrillation, Cytokine storm, Deep vein thrombosis, Haemophagocytic lymphohistiocytosis, Nervous system disorder, Platelet count
SMQs:, Supraventricular tachyarrhythmias (narrow), Embolic and thrombotic events, venous (narrow), Thrombophlebitis (broad), Hypersensitivity (broad), Tumour lysis syndrome (broad),

Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
  Date died: 2021-06-12
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: Platelets; Result Unstructured Data: Test Result:15,000
CDC Split Type: USPFIZER INC2021879891

Write-up: Neurologic damage (Shaking/memory/speech/motorskills); DVT; Afib; Secondary hlh; Cytokine Storm; This is a spontaneous report from a Pfizer-sponsored program. A female patient of an unspecified age received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on 29Mar2021 as single dose for covid-19 immunization. The patient medical history was not reported. The patient''s concomitant medications were not reported. Historical vaccine included first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection; Lot number: unknown), via an unspecified route of administration on unspecified date as single dose for COVID-19 immunization. The patient experienced neurologic damage (Shaking/memory/speech/motorskills), DVT, Afib, secondary hlh, cytokine storm on 2021. The patient was hospitalized for neurologic damage (Shaking/memory/speech/motorskills), dvt, Afib, Secondary hlh and cytokine storm from 12Apr2021 to an unknown date. The patient underwent lab tests and procedures which included platelet count: 15,000. The patient died on 12Jun2021. It was not reported if an autopsy was performed. The outcome of all the events was Fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender''s Comments: Based on temporal association, the causal relationship between BNT162B2 and the events 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.,Linked Report(s) : US-PFIZER INC-2021879892 same source, different reporter/ patient; Reported Cause(s) of Death: The Pfizer shots killed my beautiful mother

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VAERS ID: 1486322 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-07-20

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebrovascular accident
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021856673

Write-up: stroke; This is a spontaneous report from a contactable other hcp. The other hcp reported for both the husband and wife, this is the case for wife. A female patient of an unspecified age received bnt162b2 (COVID-19 VACCINE - MANUFACTURER UNKNOWN), via an unspecified route of administration on unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient experienced stroke on unspecified date and died. It was not reported if an autopsy was performed. Information about batch/lot number has been requested.; Sender"s Comments: Based on known drug safety profile, there is reasonable possibility of causal association between the event Cerebrovascular accident and the suspect drug 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.,Linked Report(s) : US-PFIZER INC-2021856461 Same reporter/ drug/event, different patients; Reported Cause(s) of Death: stroke

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VAERS ID: 1489530 (history)
Form: Version 2.0
Age: 40.0

Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210732113

Write-up: PASSED AWAY; This spontaneous report received from a consumer via a company representative via social media concerned a 5 decade old male. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported, expiry date: unknown) dose was not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient passed away due to vaccination. The cause of death was unknown. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210732113-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1489537 (history)
Form: Version 2.0
Age:
Sex: Male

Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210733844

Write-up: DEATH; This spontaneous report received from a consumer concerned a 5 decade old male of unknown race and ethnicity. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported, UNKNOWN expiry) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). It was unknown whether the Autopsy was performed.; Sender''s Comments: V0: 20210733844-covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s); Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1489604 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown

Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-21

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Death; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Not Provided Concomitant product was not reported. Treatment was not reported. Company Comment : Very limited information regarding these events has been provided at this time. Further information has been requested.; Sender''s Comments: Very limited information regarding these events has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1489619 (history)
Form: Version 2.0
Age: 81.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-07-21

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-07-16
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Died of covid; This spontaneous case was reported by a consumer and describes the occurrence of COVID-19 (Died of covid) in an 81-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced COVID-19 (Died of covid) (seriousness criterion death). The patient died on 16-Jul-2021. It is unknown if an autopsy was performed. No concomitant medications were reported. No treatment information was reported. Very limited information regarding this event has been provided at this time. Further information has been requested. Further information is not expected.; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested. Further information is not expected.

VAERS ID: 1490628 (history)
Form: Version 2.0
Age: 81.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-21

Vaccination / Manufacturer	Lot /	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Condition aggravated, Death, Diabetes mellitus, Functional gastrointestinal disorder, Hypertension, Recurrent cancer, Throat cancer
SMQs:, Hyperglycaemia/new onset diabetes mellitus (narrow), Neuroleptic malignant syndrome (broad), Oropharyngeal neoplasms (narrow), Gastrointestinal nonspecific inflammation (narrow), Hypertension (narrow), Non-haematological malignant tumours (narrow), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-04-11
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Type 2 diabetes, high blood pressure inactive remission throat cancer.
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: Bowel issues. After vaccination all of her medical conditions became worse. (type 2 diabetes, high blood pressure, and throat cancer was no longer in remission.)

VAERS ID: 1493247 (history)
Form: Version 2.0
Age:
Sex: Female
Location: South Carolina
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-22

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

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Antibody test, COVID-19, Cough, Drug ineffective, Dyspnoea, SARS-CoV-2 test
SMQs:, Anaphylactic reaction (broad), Lack of efficacy/effect (narrow), 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? Yes
  Date died: 2021-07-10
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: Bypass surgery (She had bypass 19 years ago); COPD
Allergies:
Diagnostic Lab Data: Test Name: Antibody Test; Result Unstructured Data: Test Result:no antibody at all; Test Date: 2021; Test Name: Covid-19; Test Result: Positive ; Comments: She caught Covid and died.
CDC Split Type: USPFIZER INC2021881403

Write-up: My mother in law received the vaccine 4-5 months ago. She caught Covid and died; My mother in law received the vaccine 4-5 months ago. She caught Covid and died; Just couldn"t breathe; A little cough; This is a spontaneous report received from a contactable consumer via Pfizer sponsored program. The caller is calling on behalf of her deceased Mother-in-law. A 73-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Solution for injection, Batch/Lot number was not reported, Expiration Date: Unknown), via an unspecified route of administration on an unspecified date as dose number unknown, single for COVID-19 immunization. Medical history included COPD from an unknown date and unknown if ongoing, Bypass (She had bypass 19 years ago) from an unknown date and unknown if ongoing. The reporter stated that " I don"t know what else was she taking." The patient"s concomitant medications were not reported. Reporter wants to know if Pfizer was giving a placebo or the actual vaccine. She believes her mother-in-law had received a placebo vaccine. My mother-in-law received the vaccine 4-5 months ago. She caught Covid and died (2021). The doctor confirmed she did not have any antibodies. We all had been exposed to her. We are currently in quarantine. My father-in-law is 80 years old and was nervous. He was wondering why did he take the vaccine if it did not protect his wife or him. Reported doctor had basically tested her and they had found no antibodies. At the moment she was nervous because her father-in-law lived with them and they were both quarantined and he was 80 years old. Reporter stated they said she had no antibodies like she never got the vaccine. Reporter stated I do not know the name of antibody Test, but the hospital did it and they said she had no antibody at all. Date of Death: Reporter stated, I am not sure my father-in-law did knows exactly when they got vaccinated but it was 3 or 5 months ago. Reporter stated, "No, she went to the hospital on Thursday evening, and she had been on the ventilator like 3 hours later and she died on Saturday morning 2 O"clock. She had no fever, she had a little cough not major stuff, it was like flu the fever and usual cough she did not had any of that. She had no flu, she just couldn"t breathe (2021). Anatomical Site of Administration: Reporter stated, I just know that she received it 4 months ago. The patient underwent lab tests and procedures which included antibody test: no antibody at all, Covid-19: positive on 2021 (She caught Covid and died). Therapeutic measures were taken as a result

of my mother in law received the vaccine 4-5 months ago. she caught covid and died, just couldn"t breathe. The patient died on 10Jul2021. An autopsy was not performed. The outcome of events just couldn"t breathe and cough was unknown, while other events was fatal. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: My mother in law received the vaccine 4-5 months ago. She caught Covid and died

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VAERS ID: 1493382 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-22

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Knows of some people and friends that have died after receiving Moderna"s COVID-19 vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Knows of some people and friends that have died after receiving Moderna"s COVID-19 vaccine) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Very limited information regarding this event has been provided at this time. No further follow-up information is expected. Reporter did not allow further contact; Sender"s Comments: Very limited information

regarding this event has been provided at this time. No further follow-up information is expected.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1497584 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-23

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Haemorrhagic stroke, Immune thrombocytopenia
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Systemic lupus erythematosus (broad), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021898328

Write-up: refractory ITP; hemorrhagic stroke; This is a literature report from the 2021, Patient died after receiving COVID-19 vaccine under investigation. Full Publication has been requested. 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 medical history was not reported. The patient''s concomitant medications were not reported. The patient experienced refractory ITP and hemorrhagic stroke, on an unspecified date. The patient died several weeks after receiving the Pfizer-BioNTech vaccine due to refractory ITP and hemorrhagic stroke. The

patient died on an unspecified date. It was not reported if an autopsy was performed. Information about lot/batch number cannot be obtained.; Sender''s Comments: As there is limited information in the case provided, the causal association between the events Immune thrombocytopenia, hemorrhagic stroke and the suspect drug BNT162B2 cannot be excluded. The case will be reassessed once new information is available. The impact of this report on the benefit-risk profile of the Pfizer product and on the conduct of the study 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.; Reported Cause(s) of Death: refractory ITP; hemorrhagic stroke

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VAERS ID: 1497692 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-23

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident, Death
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210741198

Write-up: DEATH; STROKE; This spontaneous report received from other health professional via a company representative from social media concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. No past medical history or concurrent

conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, frequency 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced stroke. On an unspecified date, the patient died from unknown cause of death. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of stroke was not reported. This report was serious (Death, and Other Medically Important Condition). This case, from the same reporter is linked to 20210740986 and 20210740726.; Sender"s Comments: V0: This social media received cased concerns a patient of unspecified age, gender, and ethnicity who had a "stroke" and died from an unspecified cause on unspecified date after receiving. No other pertinent details reported. Information is very limited in this case. The information available precludes a complete and meaningful assessment; hence, the causality is considered unclassifiable due to insufficient information.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1500569 (history)
Form: Version 2.0
Age: 70.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-24

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

Administered by: Other **Purchased by:** ?
Symptoms: Adverse event, COVID-19
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Diabetes
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210743441

Write-up: SUSPECTED COVID-19 INFECTION; COMPLICATION FROM VIRUS; This spontaneous report received from a patient via a company representative via social media concerned an 8 decade old male. The patient''s height, and weight were not reported. The patient''s concurrent conditions included diabetes. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient died on sunday from suspected covid-19 infection after he had been hospitalized (date unspecified) due to complications from the virus. It was reported that the patient had received vaccine, but he contracted the virus five days after the his shot before he was fully protected. it was reported that the patient was vaccinated, but he was vaccinated too late. It was not effective yet. It was stated that the patient wanted to get vaccinated, but did not get around to it for a while, and unfortunately did not get around to it until it was too late. The patient was not connected to any outbreak and he was exposed to the virus by a community member. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of suspected covid-19 infection on an unspecified date, and the outcome of complication from virus was not reported. This report was serious (Death, and Hospitalization Caused / Prolonged).; Sender''s Comments: V0- 20210743441-Covid-19 vaccine ad26.cov2.S- Suspected Covid-19 infection , Complication from virus. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: COVID INFECTION AFTER VACCINATION/COMPLICATION FROM VIRUS

VAERS ID: 1500621 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021868584

Write-up: Died in his sleep; This is a follow-up spontaneous report from a Pfizer Sponsored Program. A contactable consumer reported a 16-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date (at unknown age) as single dose for COVID-19 immunization. The patient medical history and concomitant medications were not reported. Caller stated that a 16-year-old boy died in his sleep after receiving his Covid-19 vaccine per Caller. Caller thinks it was the Pfizer Covid-19 vaccine. States that people died and adverse events happen after receiving the Pfizer Covid-19 vaccine. Regarding the 16 -ear-old male, caller is asking how could that happen to a young man that seems to be healthy? The patient died on an unspecified date. It was unknown if an autopsy was performed. information on the lot/batch number has been requested.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-2021868538 same reporter/ drug, different AE/ patient; Reported Cause(s) of Death: Died in his sleep

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VAERS ID: 1504860 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-27

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, SARS-CoV-2 test, Suspected COVID-19
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: over 80 years old patients had underlying health issues.
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Result Unstructured Data: Positive
CDC Split Type: USJNJFOC20210741034

Write-up: DEATH; SUSPECTED COVID 19 INFECTION; This spontaneous report received from a company representative via Manufacturer on 16-JUL-2021 concerned multiple patients. Initial information was processed along with the additional information received on 21-JUL-2021. The patient''s height, and weight were not reported. Two patient''s over 80 years old had underlying health issues, and pre-existing medical conditions of other 8 patient''s were not reported. The patient''s received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose, 1 in total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, two patient''s over 80 years old died from an unknown cause after being hospitalized, and eight patients also required hospitalization (hospitalization date unspecified). All patient''s had contracted the virus more than two weeks after being fully vaccinated. The cases are confirmed with a positive test COVID-19 (coronavirus disease) (confirmatory test was not reported) (suspected covid 19 infection). Laboratory data (dates unspecified) included: COVID-19 virus test (NR: not provided) Positive. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of death on an unspecified date, and the outcome of suspected covid 19 infection was not reported. This report was serious (Death, and Hospitalization Caused / Prolonged). This report was associated with product quality complaint, 90000186727.; Sender''s Comments: V0 20210741034-COVID-19 VACCINE AD26.COV2.S-Death and suspected covid 19 infection. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1504869 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-27

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210746676

Write-up: DEATH; This spontaneous report received from a patient concerned multiple patients. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin were not reported)1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210746676-covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1505007 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-27

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

Administered by: Other **Purchased by:** ?
Symptoms: Adverse event, Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: The patient was healthy.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210745184

Write-up: DEATH; FEEL SIDE EFFECTS/SYMPTOMS; This spontaneous report received from a consumer concerned a 49 year old male an unspecified race and ethnic origin. The patient''s height, and weight were not reported. The patient was healthy. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown; expiry date: Unknown) dose, start therapy date were not reported 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date (1 day after receiving vaccine), the patient experienced side effects/symptoms. Four day later, he passed away. It was not reported, if the autopsy was performed or not. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The patient died of death on an unspecified date, and the outcome of feel side effects/symptoms was not reported. This report was serious (Death). This case is linked to 20210608311 (same reporter).; Sender''s Comments: V0 - 20210745184-Covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1507809 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-28

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210747770

Write-up: DEATH; This spontaneous report received from a consumer concerned multiple patients of unspecified age, sex, race and ethnic origin. The weight, height, and medical history of patients were not reported. The patients received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patients died from unknown cause of death. The autopsy details were not provided. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210747770-Covid-19 vaccine ad26.cov2.s -Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1507874 (history)
Form: Version 2.0
Age: 91.0
Sex: Female
Location: Iowa
Vaccinated: 2021-02-01
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Cellulitis, Dementia, Dementia Alzheimer's type, Hypotension, Memory impairment
SMQs:, Anaphylactic reaction (broad), Neuroleptic malignant syndrome (broad), Dementia (narrow), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Depression (excl suicide and self injury) (broad), Infective pneumonia (broad), Dehydration (broad), Hypokalaemia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Dementia
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: blood pressure became very low; experienced cellulitis in her legs; Diagnosed with COVID a week later; she couldn''t even remember anybody; acute Alzheimers; had mild dementia but towards the end it seemed to have exacerbated; This spontaneous case was reported by a consumer and describes the occurrence of DEMENTIA ALZHEIMER''S TYPE (acute Alzheimers), DEMENTIA (had mild dementia but towards the end it seemed to have exacerbated), HYPOTENSION (blood pressure became very low) and CELLULITIS (experienced cellulitis in her legs) in a 91-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Dementia. In February 2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. In February 2021, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced DEMENTIA ALZHEIMER''S TYPE (acute Alzheimers) (seriousness criteria death and medically significant), DEMENTIA (had mild dementia but towards the end it seemed to have exacerbated) (seriousness criterion medically significant), HYPOTENSION (blood pressure became very low) (seriousness criterion hospitalization), CELLULITIS (experienced cellulitis in her legs) (seriousness criterion medically significant), COVID-19 (Diagnosed with COVID a week later) and MEMORY IMPAIRMENT (she couldn''t even remember anybody). The patient died on an unknown date. The reported cause of death was death was acute alzheimers. It is unknown if an autopsy was performed. At the time of death, DEMENTIA (had mild dementia but towards the end it seemed to have exacerbated), CELLULITIS (experienced cellulitis in her legs), COVID-19 (Diagnosed with COVID a week later) and MEMORY IMPAIRMENT (she couldn''t even remember anybody) outcome was unknown and HYPOTENSION (blood pressure became very low) had resolved. Treatment included unspecified pressors. No concomitant medications were reported. Patient received the 1st dose of the Moderna COVID-19 vaccine in FEB-2021. Caller did not know the exact dates of administration of the two doses. Patient was then diagnosed with COVID a week later. Patient did not have much of a reaction to the 1st dose or any symptoms from COVID. She then received her 2nd dose of the Moderna COVID-19 vaccine in middle of FEB-2021, per the caller. In the end of APR-2021 or beginning of MAY-2021, patient experienced cellulitis in her legs. Her blood pressure became very low and she was hospitalized for treatment. She was put on pressors. She recovered from it and was sent back to her assisted living facility. However, on 25-MAY-2021, she passed away at the assisted living due to acute Alzheimers. Patient had mild dementia but towards the end it seemed to have exacerbated, where she couldn''t even remember anybody. Her diagnosis at the time of death was acute Alzheimers. This case was linked to MOD-2021-034532 (Patient Link).; Sender''s Comments: This is a 91-year-old female patient who received mRNA-1273 (COVID 19 Vaccine Moderna) (batch unknown) who experienced exacerbation of her DEMENTIA, ALZHEIMER''S DEMENTIA, HYPOTENSION and CELLULITIS after her second dose of vaccine, leading to her death from ACUTE ALZHEIMERS. Advanced senior age also

could be attributed to the development of the events. It is unlikely, that the events are related to the vaccine based on long standing illness of DEMENTIA. . Very limited information has been reported at this time. Further information is expected.; Reported Cause(s) of Death: death was acute Alzheimers

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VAERS ID: 1507888 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100913512

Write-up: Caller states that 5 deaths, from that vaccine was reported; This is a spontaneous report from a contactable consumer. This is the 1st of 5 reports. A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, formulation: solution for injection, Lot number: unknown) via an unspecified route of administration on an unspecified date as dose number unknown, single for Covid-19 immunization. The patients medical history and concomitant medications were not reported. On an unspecified date, Consumer stated that 5 deaths, from that vaccine was reported and over 50% of people reported a debilitating disease, or cannot function, and have thousands of dollars in medical bills. The outcome of event was fatal. Autopsy was unknown. Information on the lot/ batch number has been requested. ; Reported Cause(s) of Death: Caller states that 5 deaths, from

that vaccine was reported

VAERS ID: 1507903 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100944572

Write-up: Death; This is a spontaneous report from a non-contactable consumer. This is one of 5 reports A patient of unspecified age and gender received bnt162b2 (BNT162B2) dose number unknown, via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient died on an unknown date from an unknown cause. It was not reported if an autopsy was performed. The reporter stated that the vaccine was kept in ice box for like 5 hours. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100944710 same reporter/drug/event, different patient;US-PFIZER INC-202100944708 same reporter/drug/event, different patient;US-PFIZER INC-202100944711 same reporter/drug/event, different patient;US-PFIZER INC-202100944709 same

reporter/drug/event, different patient; Reported Cause(s) of Death: Death

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VAERS ID: 1507904 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100944708

Write-up: Death; This is a spontaneous report from a non-contactable consumer. This is one of 5 reports A patient of unspecified age and gender received bnt162b2 (BNT162B2) dose number unknown, via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient died on an unknown date from an unknown cause. It was not reported if an autopsy was performed. The reporter stated that the vaccine was kept in ice box for like 5 hours. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100944572 same reporter/drug/event, different patient; Reported Cause(s) of Death: Death

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VAERS ID: 1509030 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC2021889151

Write-up: passed away; 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 a single dose for covid-19 immunisation. Medical history and concomitant medications were unknown. No patient identifiers were provided but the reporter has firsthand knowledge of the patient and was reporting on a specific patient. The reporter felt people will get healthier with their own immune systems and that Covid was a cover up. She knew people that have died after getting vaccinated and the CDC had changed their guidelines many times now. She read if 50% were vaccinated and the other half combined with the virus but now that had changed to 70%. The body had been proven for antibodies to work and it''s no different than the flu shot. The reporter had a 16 year old girl come to her door today letting her know that her neighbor passed away and she asked the 16-year-old if she could hug her and the little girl said "yes I''ve been vaccinated". It was unknown if autopsy was performed. Outcome of the event was fatal. The lot number for the vaccine [BNT162B2] was not provided and will be requested during follow up.; Reported Cause(s) of Death: passed away

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VAERS ID: 1511553 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210748471

Write-up: DEATH; SUSPECTED CLINICAL VACCINATION FAILURE; SUSPECTED COVID-19 INFECTION; This spontaneous report received via social media (news article) and concerned 26 multiple patients. Initial information was processed along with the additional information received from Regulatory Authority on 23-JUL-2021 and 27-JUL-2021. The patient"s weight, height, and medical history were not reported. The patient"s received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose was not reported, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patients had a breakthrough infection after being vaccinated (suspected clinical vaccination failure and suspected covid-19 infection). It was reported that it was only 25 to 54 days after that last dose of the vaccine when the patients got sick. It was also reported that, on an unspecified date vaccinated patients were hospitalized and four experienced death from unknown cause of death. The number of days hospitalization was not reported. It was unknown if an autopsy was performed to the patients. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died on an unspecified date, and the outcome of suspected clinical vaccination failure and suspected covid-19 infection was not reported. This report was serious (Death, and Hospitalization Caused / Prolonged). This report was

associated with product quality complaint number: 90000186897.; Sender''s Comments: V0:20210748471-COVID-19 VACCINE AD26.COV2.S-Death, suspected COVID-19 infection . This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).20210748471-COVID-19 VACCINE AD26.COV2.S-Suspected clinical vaccination failure. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1511577 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-29

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210756409

Write-up: DIED 24 HOURS LATER AFTER TAKING VACCINE; This spontaneous report received from a company representative via Pfizer. via social media was received on 23-JUL-2021 and concerned a 21 year old of an unspecified sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, start therapy date was not

reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient died 24 hours later after taking vaccine. On an unspecified date, the patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: 20210756409-Covid-19 vaccine ad26.cov2.s -died 24 hours later after taking vaccine. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1511611 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-29

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Death after vaccination; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death after vaccination) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. An autopsy was

not performed. Concomitant medication and treatment information were not reported. Company Comment: Very limited information regarding this event has been provided at this time. Further information is not expected (Follow up contact denied) Reporter did not allow further contact; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information is not expected (Follow up contact denied); Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1511613 (history)
Form: Version 2.0
Age: 24.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-29

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Asthenia, Dyspnoea, Hypokinesia, Lung disorder, Neuromyelitis optica spectrum disorder
SMQs:, Anaphylactic reaction (broad), Parkinson-like events (broad), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Optic nerve disorders (narrow), Cardiomyopathy (broad), Demyelination (narrow), Hypotonic-hyporesponsive episode (broad), Immune-mediated/autoimmune disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-07-08
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: USMODERNATX, INC.MOD20212

Write-up: weakness; difficulty breathing; unable to move his legs; Lung problem; neuromyletis optica; This spontaneous case was reported by a consumer and describes the occurrence of NEUROMYELITIS OPTICA SPECTRUM DISORDER (neuromyletis optica), ASTHENIA (weakness), DYSPNOEA (difficulty breathing), HYPOKINESIA (unable to move

his legs) and LUNG DISORDER (Lung problem) in a 24-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced NEUROMYELITIS OPTICA SPECTRUM DISORDER (neuromyletis optica) (seriousness criteria death, hospitalization and medically significant), ASTHENIA (weakness) (seriousness criteria death and hospitalization), DYSPNOEA (difficulty breathing) (seriousness criteria death and hospitalization), HYPOKINESIA (unable to move his legs) (seriousness criteria death and hospitalization) and LUNG DISORDER (Lung problem) (seriousness criteria death and hospitalization). The patient was hospitalized on 09-Jun-2021 due to ASTHENIA, DYSPNOEA, HYPOKINESIA, LUNG DISORDER and NEUROMYELITIS OPTICA SPECTRUM DISORDER. The patient died on 08-Jul-2021. The reported cause of death was Neuromyelitis optica. It is unknown if an autopsy was performed. No concomitant medication was reported. The patient was placed on ventilator and had a tracheotomy tube. No treatment information was provided. No laboratory data was provided. Very limited information regarding the events has been provided at this time. Reporter did not allow further contact; Sender"s Comments: Very limited information regarding the events has been provided at this time.; Reported Cause(s) of Death: Neuromyelitis optica

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VAERS ID: 1511620 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-29

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebral haemorrhage, Neoplasm malignant
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Non-haematological malignant tumours (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100911615

Write-up: Caller states that the woman in the study had brain hemorrhaging and a cancerous growth, after the vaccine.; Caller states that the woman in the study had brain hemorrhaging and a cancerous growth, after the vaccine.; This is a spontaneous report from a Pfizer-sponsored program Support reported by a contactable consumer or other non-health care professional. A female patient of an unspecified age received unknown dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number and Expiration date not reported), via an unspecified route of administration in an unspecified anatomical location on an unspecified date as a single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. Reporter stated that cardiologist gave him a study which reported a woman, who had the vaccine, died on the operating table, after her brain was haemorrhaging and they found a cancerous growth. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information about Lot/ Batch number have been requested.; Reported Cause(s) of Death: after her brain was hemorrhaging; and they found a cancerous growth

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VAERS ID: 1517882 (history)
Form: Version 2.0
Age:
Sex: Female
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Other **Purchased by:** ?
Symptoms: Aortic valve incompetence, Blood bicarbonate, Blood creatine phosphokinase, Blood creatinine, Blood culture, Blood lactic acid, Blood pressure measurement, Body mass index, C-reactive protein, Death, Echocardiogram, Electrocardiogram, Heart rate, Incorrect dose administered, Myocarditis, Off label use, Oxygen saturation, PCO2, PO2, Procalcitonin, Pulmonary arterial wedge pressure, Renal impairment, Respiratory failure, Respiratory rate, Septic shock, Troponin, pH body fluid
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Anaphylactic reaction (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 (narrow), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Acute central respiratory depression (narrow), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Hypersensitivity (broad), Tumour lysis syndrome (broad), Respiratory failure (narrow), Medication errors (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Hypokalaemia (broad), Sepsis (narrow), Opportunistic

infections (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 8 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Multiple sclerosis
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data: Test Name: Heart rate; Result Unstructured Data: 145 bpm; Test Name: Heart rate; Result Unstructured Data: 125 BPM; Test Name: Blood pressure; Result Unstructured Data: 117/70 mmHg; Test Name: Respiratory rate; Result Unstructured Data: 39; Test Name: Body mass index; Result Unstructured Data: 27.5; Test Name: Electrocardiogram; Result Unstructured Data: sinus tachycardia; Test Name: Blood culture; Result Unstructured Data: Negative for all viruses; Test Name: Procalcitonin; Result Unstructured Data: 185.71 ng/mL; Test Name: Troponin; Result Unstructured Data: 1.260-2.050 ng/mL; Test Name: Transthoracic echocardiogram; Result Unstructured Data: 2+ aortic regurgitation and diffuse left ventricular hypokinesis; Test Name: Echocardiogram; Result Unstructured Data: Diffuse left ventricular hypokinesis with severely reduced contraction; Test Name: Pulmonary arterial wedge pressure; Result Unstructured Data: 14 mmHg; Test Name: Creatinine; Result Unstructured Data: 1.21 mg/dL; Test Name: Bicarbonate; Result Unstructured Data: 16 mmol/L; Test Name: Troponin; Result Unstructured Data: <0.010 ng/mL; Comments: reference range =0010 ng/mL; Test Name: Creatine phosphokinase; Result Unstructured Data: 53 U/L; Test Name: Procalcitonin; Result Unstructured Data: 0.07 ng/mL; Test Name: C-reactive protein; Result Unstructured Data: 7.2 mg/L; Test Name: pH; Result Unstructured Data: 7.02; Test Name: Partial pressure CO2; Result Unstructured Data: 94 mmHg; Test Name: PaO2; Result Unstructured Data: 27 mmHg; Test Name: Lactate; Result Unstructured Data: 8.3 mmol/L; Test Name: Creatinine; Result Unstructured Data: worsening; Test Name: Oxygen saturation; Result Unstructured Data: 75 %
CDC Split Type: USJNJFOC20210754113

Write-up: VACCINE INDUCED MYOCARDITIS (VIRAL MYOCARDITIS); ACUTE HYPOXIC HYPERCAPNIC RESPIRATORY FAILURE; SEPTIC SHOCK; DECLINE IN RENAL FUNCTION; 2+ AORTIC REGURGITATION; PATIENT ADMINISTERED WITH 2 DOSES OF VACCINE; OFF LABEL USE; DEATH; This spontaneous report was received from literature: Case report with literature review. This report concerned a 70 year old female. The patient''s height, and weight were not reported. The patient''s concurrent conditions included multiple sclerosis. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown), frequency 2 total doses, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, after two days post second dose (inappropriate dose of vaccine administered and off label use) the patient experienced developed dyspnea at home and eventually required an ambulance for hospital transfer. The vital signs on arrival included a heart rate of 145 bpm, a 75% oxygen saturation level on room air, a blood pressure of 117/70

mmHg, a respiratory rate of 39, and a BMI of 27.5. The electrocardiogram (ECG) on admission revealed sinus tachycardia with a heart rate of 125bpm and T-wave inversions in leads V4-V6 without any ST-segment change. The patient arrived at the emergency department in severe respiratory distress that warranted immediate intubation. She was admitted to the intensive care unit (ICU) with the provisional diagnoses of acute hypoxic hypercapnic respiratory failure and septic shock. The laboratory screening and blood culture proved negative for all viruses, Mycoplasma pneumonia, and chlamydophila pneumonia. A repeat investigation revealed marked elevations in procalcitonin [185.71(ng/mL)] and troponin [1.260-2.050 ng/mL] levels on the second day of admission. The patient required multiple vasopressors to maintain the mean arterial pressure above 65 mmHg. The transthoracic echocardiogram on admission revealed 2+ aortic regurgitation and diffuse left ventricular hypokinesis with severely decreased left ventricular ejection fraction (10%). A repeat echocardiogram with contrast medium showed diffuse left ventricular hypokinesis with diagnostic monitoring via Swanz-Ganz catheter revealed a pulmonary wedge pressure (PWP) of 14mmHg. The patient continued receiving vasopressors and antibiotic therapy, while her renal function deterioration since admission warranted the prompt administration of renal replacement therapy. Further decline in renal function was marked by oliguria and worsening of creatinine levels. The patient declined cardiac catheterization and remained. On an unspecified date the patient was hospitalized and it was for 8 days. On unspecified date patient died with vaccine induced myocarditis (viral myocarditis) on eighth day of admission to hospital. The exact cause of death was not reported and it was unknown whether autopsy was performed. The other laboratory data includes, Creatinine (NR: 0.05 - 1.20) 1.21 mg/dL, Bicarbonate (NR: 22 - 29) 16 mmol/L, Creatine phosphokinase (NR: 20 - 190) 53 U/L, Procalcitonin (NR: 0.02 - 0.10) 0.07 ng/mL, C-reactive protein (NR: 0 - 3.00) 7.2 mg/L, pH (NR: 7.35 - 7.45) 7.02 (units unspecified), Partial pressure CO2 (NR: 35 - 48) 94 mmHg, Lactate (NR: 0.6 - 1.4) 8.3 mmol/L, PaO2 (NR: 83 - 108) 27 mmHg, The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of vaccine induced myocarditis (viral myocarditis), acute hypoxic hypercapnic respiratory failure, septic shock, decline in renal function, 2+ aortic regurgitation, patient administered with 2 doses of vaccine, off label use was not reported. This report was serious (Death, Life Threatening, and Hospitalization Caused / Prolonged).; Sender"s Comments: V0: 20210754113-covid-19 vaccine ad26.cov2.s - Death, vaccine induced myocarditis, acute hypoxic hypercapnic respiratory failure, septic shock, decline in renal function, 2+ aortic regurgitation. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1517885 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Massachusetts
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

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

Symptoms: Death, SARS-CoV-2 test, Suspected COVID-19, Vaccine breakthrough infection
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: BNT 162; MRNA 1273
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Result Unstructured Data: Positive
CDC Split Type: USJNJFOC20210756412

Write-up: DEATH; BREAKTHROUGH COVID CASES; SUSPECTED COVID 19 INFECTION; This spontaneous report received from a patient by other company (Pfizer Inc.) received on 23-JUL-2021 and concerned multiple patients. No past medical history or concurrent conditions were reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. Concomitant medications included bnt 162, and mrna 1273. It was reported that there were 4450 breakthrough Covid cases. Among 4450 cases, 247 patients were hospitalized for suspected Covid 19 infection and 79 patients died among the population who were fully vaccinated by Pfizer or Moderna vaccine both doses or Janssen vaccine one dose. Laboratory data (dates unspecified) included: COVID-19 virus test (NR: not provided) Positive. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of death on an unspecified date, and the outcome of breakthrough covid cases and suspected covid 19 infection was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: V0:20210756412-covid-19 vaccine ad26.cov2.s-death,breakthrough covid cases. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210756412-covid-19 vaccine ad26.cov2.s -suspected covid 19 infection. This event(s) is considered not related. The event(s) is deemed to be scientifically implausible, i.e., there is scientific evidence against a drug/event relationship; AND there is no known class effect.; Reported Cause(s) of Death: CONFIRMED COVID-19 INFECTION

VAERS ID: 1517971 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100909825

Write-up: Reporter had a friend die from the covid vaccine.; This is a spontaneous report from a non-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 (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Reporter had a friend die from the covid vaccine. Outcome of event was fatal.The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Reporter had a friend die from the covid vaccine.

VAERS ID: 1517987 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

Vaccination / Manufacturer	Lot /	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100912595

Write-up: died from the vaccine; This is a spontaneous report from a contactable consumer reporting same event under the same suspect product for 4 patients. This is one of four 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 (Batch/Lot number was not reported) 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 died from the vaccine on an unspecified date. The reporter stated in his neighborhood 4 people died from the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. Follow-up attempts have been completed and no further information is expected. Information about lot/batch number cannot be obtained.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202100970595 same reporter, event and drug, different patient;US-PFIZER INC-202100970596 same reporter/drug/event, different pt;US-PFIZER INC-202100970597 same reporter, event and drug, different patient; Reported Cause(s) of Death: died from the vaccine

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VAERS ID: 1518030 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Sudden death
SMQs:, Torsade de pointes/QT prolongation (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100922421

Write-up: sudden death of a 13-year-old (place)child who had a second dose of Pfizer vaccine; This is a spontaneous report from a non-contactable consumer (columnist). A 13-year-old male patient received second dose of BNT162b2 (FIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. It was reported that remained strangely silent over the sudden death of a 13-year-old child who had a second dose of Pfizer vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible, information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Sudden death

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VAERS ID: 1518133 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100944709

Write-up: 5 death happen; This is a spontaneous report from a non-contactable consumer. This consumer reporting same event under the same suspect product for 5 patients. This is 3rd of 5 reports. A patient of an unknown age and gender received 2nd dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) at single dose via an unknown route on an unknown date for Covid-19 immunization. Medical history and concomitant drug were not provided. Patient died after Pfizer vaccine. It was further reported that couple of deaths occurred after Pfizer vaccine, some of them immediately, some in few hours. The reporter stated that in one village with 2000 population, 5 death happen and some after the 2nd vaccine and one of them immediately after vaccine, The vaccine was kept in ice box for like 5 hours, other village it just about 3, another village or 2. The reporter stated if the vaccine was carried by just ice for like 5 hour, may be this was the problem. Outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202100944572 same reporter/drug/event, different patient; Reported Cause(s) of Death: 5 death happen

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VAERS ID: 1518134 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100944710

Write-up: deaths occurred after Pfizer vaccine; This is a spontaneous report from a non-contactable consumer. This consumer reported similar events for 5 patients, this is the forth report. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via unspecified route single dose for COVID-19 immunization on an unspecified date. The patient medical history was not reported. The patient''s concomitant medications were not reported. Patient died after Pfizer vaccine. It was reported that couple of deaths occurred after Pfizer vaccine, some of them immediately, some in few hours. The reporter stated that in one village with 2000 population, 5 death happen and some after the 2nd vaccine and one of them immediately after vaccine, The vaccine was kept in ice box for like 5 hours, other village it just about 3, another village or 2. The reporter asked if the vaccine was carried by just ice for like 5 hour, may be this was the problem. Outcome of the event was fatal. The patient died on an unspecified date. It was unknown if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100944572 same reporter/drug/event, different patient; Reported Cause(s) of Death: deaths occurred after Pfizer vaccine

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VAERS ID: 1518135 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100944711

Write-up: Couple of deaths occurred after Pfizer vaccine, some of them immediately, some in few hour/5 death happen and some after the 2nd vaccine and one of them immediately after vaccine; This is a spontaneous report from a non-contactable consumer. This consumer reported similar events for 5 patients, this is the 5th report. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient died on an unspecified date. Reporter wanted to ask about the question about moving the vaccine in packets of ice for like 2or 3 or 4 hour. Will it make it less effective or may be bad effect? Reporter stated, the question was about couple of deaths occurred after Pfizer vaccine, some of them immediately, some in few hour. Their question was about the vaccine that was carried by just ice for like 5 hour, may be this was the problem. Reporter stated, everything is fine you know in one village with 2000 population, 5 death happen and some after the 2nd vaccine and one of them immediately after vaccine. The vaccine was kept in ice box for like 5 hours, other village it just about 3, another village or 2. The question is, if the ice box is enough for carrying the vaccine. It was not reported if an autopsy was performed. Seriousness criteria was resulted in death. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100944572 same reporter/drug/event, different patient; Reported Cause(s) of Death: Couple of deaths occurred after Pfizer vaccine, some of them immediately, some in few hour/5 death happen and some after the 2nd vaccine and one of them immediately after vaccine

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VAERS ID: 1518168 (history)
Form: Version 2.0

Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100963386

Write-up: Caller states that 5 deaths, from that vaccine was reported; This is a spontaneous report from a contactable consumer reporting for a patient received from a Pfizer-sponsored program. This is the 2nd of 5 reports. A patient of an unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, formulation: solution for injection, Lot number: unknown) via an unspecified route of administration on an unspecified date as DOSE NUMBER UNKOWN, SINGLE for Covid-19 immunization, age at vaccination unknown. The patients medical history and concomitant medications were not reported. Consumer stated that he did took a survey in his neighborhood, going door to door, and some people had taken the vaccine and some had not. Consumer stated that 5 deaths, from that vaccine was reported, and over 50% of people reported a debilitating disease, or cannot function, and have thousands of dollars in medical bills. The outcome of event was fatal. Autopsy was unknown. No follow-up attempts are needed; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100913512 Same reporter/drug/AE, different patient; Reported Cause(s) of Death: Consumer stated that 5 deaths, from that vaccine was reported

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VAERS ID: 1518169 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100963754

Write-up: 5 deaths, from that vaccine was reported; This is a spontaneous report from a contactable consumer received from a Pfizer-sponsored program. This is the 3rd of 5 reports. A patient of an unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, formulation: solution for injection, Lot number: unknown) via an unspecified route of administration on an unspecified date as dose number unknown, single for Covid-19 immunization. The patient''s medical history and concomitant medications were not reported. On an unspecified date, Consumer stated that 5 deaths, from that vaccine was reported and over 50% of people reported a debilitating disease, or cannot function, and have thousands of dollars in medical bills. The outcome of event was fatal. Autopsy was unknown. Information on the lot/ batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100913512 Same reporter/drug/AE, different patient; Reported Cause(s) of Death: 5 deaths, from that vaccine was reported

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VAERS ID: 1518170 (history)
Form: Version 2.0

Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100963755

Write-up: Caller states that 5 deaths, from that vaccine was reported,; This is a spontaneous report from a contactable consumer received from a Pfizer-sponsored program. This consumer reported similar events for five patients. This is the 4th of five reports. A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose number unknown, single for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. Reporter called about the COVID 19 vaccine, Pfizer. The reporter did a survey in his neighborhood, going door to door, and some people had taken the vaccine and some had not. Reporter stated that 5 deaths from that vaccine was reported, and over 50% of people reported a debilitating disease, or cannot function, and have thousands of dollars in medical bills. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202100913512 Same reporter/drug/AE, different patient; Reported Cause(s) of Death: Callers states that 5 deaths, from that vaccine was reported

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VAERS ID: 1518171 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100963756

Write-up: deaths from that vaccine; This is a spontaneous report from a contactable consumer received from a Pfizer-sponsored program. This consumer reported similar events for five patients. This is the fifth of five reports. A female patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose number unknown, single for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. Reporter called about the COVID 19 vaccine, Pfizer. He did a survey in his neighborhood, going door to door, and some people had taken the vaccine and some had not. Reporter stated that 5 deaths from that vaccine was reported, and over 50% of people reported a debilitating disease, or cannot function, and have thousands of dollars in medical bills. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100913512 Same reporter/drug/AE, different patient; Reported Cause(s) of Death: deaths from that vaccine

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VAERS ID: 1518172 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100970595

Write-up: died from the vaccine; This is a spontaneous report from a contactable consumer reporting same event under the same suspect product for 4 patients. This is one of four 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 (Batch/Lot number was not reported) 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 died from the vaccine on an unspecified date. The reporter stated in his neighborhood 4 people died from the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. Follow-up attempts have been completed and no further information is expected. Information about lot/batch number cannot be obtained.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202100912595 same reporter, event and drug, different patient; Reported Cause(s) of Death: died from the vaccine

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VAERS ID: 1518173 (history)
Form: Version 2.0

Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100970596

Write-up: died from the vaccine; This is a spontaneous report from a contactable consumer reporting same event under the same suspect product for 4 patients. This is one of four 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 (Batch/Lot number was not reported) 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 died from the vaccine on an unspecified date. The reporter stated in his neighborhood 4 people died from the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. Follow-up attempts have been completed and no further information is expected. Information about lot/batch number cannot be obtained.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100912595 same reporter/drug/event, different pt; Reported Cause(s) of Death: died from the vaccine

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VAERS ID: 1518174 (history)
Form: Version 2.0
Age:

Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-07-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100970597

Write-up: died from the vaccine; This is a spontaneous report from a contactable consumer reporting same event under the same suspect product for 4 patients. This is one of four 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 (Batch/Lot number was not reported) 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 died from the vaccine on an unspecified date. The reporter stated in his neighborhood 4 people died from the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. Follow-up attempts have been completed and no further information is expected. Information about lot/batch number cannot be obtained.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202100912595 same reporter, event and drug, different patient; Reported Cause(s) of Death: died from the vaccine

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VAERS ID: 1521958 (history)
Form: Version 2.0
Age:
Sex: Female

Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-03

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Lost her mother after getting the vaccine; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Lost her mother after getting the vaccine) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No treatment information was provided. No relevant concomitant medications were reported. Very limited information regarding the events has been provided at this time; Sender''s Comments: Very limited information regarding the events has been provided at this time; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1524928 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Tennessee
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-08-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Body mass index, Body mass index abnormal, COVID-19, Drug ineffective, SARS-CoV-2 test
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: BMI; Result Unstructured Data: Test Result:greater than 40; Test Name: COVID test; Result Unstructured Data: Test Result:Died of COVID
CDC Split Type: USPFIZER INC202100958483

Write-up: Died of COVID; Died of COVID; BMI greater than 40; This is a spontaneous report received from a contactable physician communicated to a sales representative. A 23-year-old female patient received BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Solution for Injection), first dose as dose 1 single and second dose as dose 2 single, both via an unspecified route of administration on an unspecified date for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. Event took place after use of product. The patient experienced died of COVID, BMI greater than 40 on an unspecified date. The events drug ineffective and covid-19 assessed as serious (death, medically significant) and while for other was non-serious. The patient underwent lab tests and procedures which included body mass index: greater than 40, COVID test: died of covid on unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the events drug ineffective and covid-19 was reported as fatal and while for other was unknown. Reporter does not wish to be contacted for follow-up. No follow-up attempts are possible, information about lot/batch number cannot be obtained.; Sender''s Comments: (Based on plausible dose -event relationship post vaccination and no other alternate explanation the causal role of BNT162B2 cannot be excluded for the reported LOE 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.; Reported Cause(s) of Death: Died of COVID; Died of COVID

VAERS ID: 1528255 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Idaho
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-05

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Pulmonary thrombosis
SMQs:, Embolic and thrombotic events, venous (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Xeljanz
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100972826

Write-up: blood clots in her lungs; This is a spontaneous report from a contactable consumer or other non hcp from a Pfizer sponsored program. A 86-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection), dose unknown via an unspecified route of administration on an unspecified date in 2020 (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for COVID-19 immunisation; tofacitinib citrate (XELJANZ), route of administration, start and stop date, batch/lot number and dose were not reported for an unspecified indication. The patient medical history and concomitant medications were not reported. The reporter stated that his neighbor was taking Xeljanz and got the COVID 19 vaccine (unknown manufacturer) and died from blood clots in her lungs. She died on an unspecified date and the family was blaming Xeljanz. The patient was diagnosis with blood clots about a month ago. She got COVID Vaccine around Christmas 2020. She was in hospital for 5 days with blood clots. The reporter inquired if vaccine cause blood clots. Seriousness of the event was reported as death and hospitalization. The action taken in response to the event for tofacitinib citrate was not

applicable. The outcome of the event was fatal. It was not reported if an autopsy was performed. Follow-up attempts are needed. Information on the lot/batch number is expected.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1530586 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-06

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Blood grouping, Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Blood type; Result Unstructured Data: Test Result:AB Negative
CDC Split Type: USPFIZER INC202100953049

Write-up: patient died after receiving the 1st dose of the vaccine; This is a spontaneous report from a non-contactable consumer (patient''s sister) via a Pfizer sponsored program. A male patient of an unspecified age received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Batch/Lot number was not reported) via an unspecified route of administration on an unspecified date as dose 1, single for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. It was reported that the patient died after receiving the 1st dose of the vaccine on an unspecified date. The patient underwent lab tests and procedures which included blood grouping: AB negative on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is

expected.; Reported Cause(s) of Death: patient died after receiving the 1st dose of the vaccine

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VAERS ID: 1531269 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-06

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

Administered by: Other **Purchased by:** ?
Symptoms: COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-07-01
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210808115

Write-up: CONFIRMED COVID-19 INFECTION; SUSPECTED VACCINATION FAILURE; This spontaneous report received from a patient concerned a male of unspecified age, race and ethnicity unknown. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose not reported, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. It was reported that on an unspecified date, the patient experienced confirmed covid-19 infection, suspected Vaccination Failure. On an unspecified date in Jul-2021, the patient died from confirmed covid-19 infection. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of Suspected Vaccination Failure was not reported. This report was serious (Death).; Sender''s Comments: V0-20210808115-covid-19 vaccine

ad26.cov2.s-Confirmed COVID-19 infection. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210808115-covid-19 vaccine ad26.cov2.s-Suspected Vaccine failure. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS; Reported Cause(s) of Death: CONFIRMED COVID-19 INFECTION

VAERS ID: 1533998 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210810431

Write-up: DIED; BLOOD CLOT; This spontaneous report received from a consumer concerned a female of unspecified age, race and ethnicity. The patient''s weight, height and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown expiry: unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, the patient had blood clot and died. The patient died from unknown cause of death and blood clot. It was

unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. On an unspecified date, the patient died and the outcome of blood clot was fatal. This report was serious (Death).; Sender''s Comments: V0:20210810431-Covid-19 vaccine ad26.cov2.s-Death, Blood Clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH; BLOOD CLOT

VAERS ID: 1534123 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, SARS-CoV-2 antibody test
SMQs:, COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Antibody; Test Result: Negative
CDC Split Type: USPFIZER INC202100970230

Write-up: passed away; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on an unspecified date as single dose, and dose 1 via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. As reported, caller was a patient who stated he and his wife received the Pfizer COVID vaccine, and were vaccinated in Jan2021 and Feb2021. Caller was inquiring

about the antibody testing, as he and his wife had one done. Caller wanted to know why he and his wife tested negative for antibodies. Caller was concerned because his sister who had both the Pfizer COVID vaccine shots passed away and she showed not having any antibodies. The patient died on an unspecified date. An autopsy was not performed. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: passed away

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VAERS ID: 1534137 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EN6204 / 1	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100978617

Write-up: I want to report a death from a Pfizer vaccine; This is a spontaneous report from a contactable consumer (patient''s wife). A 60-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot Number: EN6207) as dose 1, single; via an unspecified route of administration on an unspecified date (Batch/Lot Number: EN6204) as dose 2, single for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient died from the Pfizer vaccine on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible. No further information is

expected.; Reported Cause(s) of Death: I want to report a death from a Pfizer vaccine

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VAERS ID: 1534138 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100978632

Write-up: They have thousands of deaths, including her mother; This is a spontaneous report from a non-contactable consumer (patient"s daughter). A female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The consumer stated they have thousands of deaths, including her mother (the patient). The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: They have thousands of deaths, including her mother

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VAERS ID: 1534143 (history)

Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100985229

Write-up: passed away, after taking Covid shots; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received bnt162b2 (BNT162B2, Lot number was not reported), via an unspecified route of administration on an unspecified date as dose 2, single for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The patient previously received the first dose of bnt162b2 on unspecified date as single dose for COVID-19 immunization. The patient passed away, after taking Covid shots on an unspecified date. Cause of death was unknown. It was unknown if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: passed away, after taking Covid shots shots

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VAERS ID: 1534144 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown

Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100985462

Write-up: My mom would still be alive today if she hadn''t been given the jab.; This is a spontaneous report from a Pfizer-sponsored program. A non-contactable consumer reported for a female patient (mom) that: A female patient of an unspecified age received bnt162b2 (BNT162B2), dose number unknown via an unspecified route of administration on an unspecified date (at an unspecified age of vaccination) (Batch/Lot number was not reported) as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient died on an unspecified date. The course of events was as follows: My mom would still be alive today if she hadn''t been given the jab. It was not reported if an autopsy was performed. No follow-up attempts are possible, information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: My mom would still be alive today if she hadn''t been given the jab.

VAERS ID: 1534147 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-08-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Illness
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100985867

Write-up: This vaccine got my grandfather sick and now he s no more; This is a spontaneous report from a Pfizer sponsored program based on information received from a Consumer about the grandfather. A male patient of an unspecified age received bnt162b2 (BNT162B2) unknown dose number, via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient became sick and died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: Sickness

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VAERS ID: 1534150 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-07

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100986086

Write-up: died after 4hours of receiving it; This is a spontaneous report from a Pfizer sponsored program received from a non-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 (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The reporter mentioned that a family member (the patient) who received the Pfizer Covid-19 vaccine died after 4hours of receiving it. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: died after 4hours of receiving it

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VAERS ID: 1542115 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-11

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis

SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210815346

Write-up: BLOOD CLOTS; This spontaneous report received from a consumer via a company representative via social media concerned multiple (3) patients. The patients` weight, height, and medical history were not reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) frequency 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patients experienced blood clots and died from blood clot. It was unknown whether the autopsy was done or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0:20210815346-Covid-19 vaccine ad26.cov2. s-Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT

VAERS ID: 1550084 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-12

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and

venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210818401

Write-up: DEATH; BLOOD CLOTS; This spontaneous report received from a consumer via social media concerned multiple patients (6 or 7) with unspecified race and ethnicity. The patient"s weight, height, and medical history were not reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patients experienced blood clots and died from unknown cause of death. It was unspecified if was an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Death, and Other Medically Important Condition). This case, from the same reporter is linked to 20210818227, 20210818107, 20210817805 and 20210818021.; Sender"s Comments: V0: 20210818401 - covid-19 vaccine ad26.cov2.s- death, Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1555771 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-14

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

Administered by: Other **Purchased by:** ?
Symptoms: Suspected COVID-19, Vaccination failure

SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-01
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: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210815127

Write-up: SUSPECTED CLINICAL VACCINATION FAILURE; SUSPECTED COVID-19; This spontaneous report received from a company representative on behalf of a health care professional concerned a male patient of unspecified age, race and ethnicity. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry: unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date after vaccination, the patient experienced suspected covid-19 (suspected clinical vaccination failure). On an unspecified date in AUG-2021 (same week of this report), the patient was hospitalized and was intubated. The duration of hospitalization was not specified. On an unspecified date in AUG-2021 (same week of this report), the patient died of covid-19. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of suspected clinical vaccination failure and suspected covid-19 on AUG-2021. This report was serious (Death, and Hospitalization Caused / Prolonged). This case was associated with the Product Quality Complaint number 90000188817.; Sender''s Comments: V0: 20210815127- JANSSEN COVID-19 VACCINE Ad26.COV2.S- Suspected COVID-19. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210815127- JANSSEN COVID-19 VACCINE Ad26.COV2.S- Suspected clinical vaccination failure. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS.; Reported Cause(s) of Death: COVID-19

VAERS ID: 1577472 (history)
Form: Version 2.0
Age:

Sex: Female
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebral haemorrhage, Thrombosis
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Haemorrhagic central nervous system vascular conditions (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210810396

Write-up: BRAIN HAEMORRHAGING; BLOOD CLOT; This spontaneous report received from a consumer who reported hearing a news report which concerned a female of unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced brain haemorrhaging and blood clot. On an unspecified date, the patient died from brain hemorrhage and blood clot. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of brain haemorrhaging and blood clot on an unspecified date. This report was serious (Death).; Sender''s Comments: V0: 20210810396- JANSSEN COVID-19 VACCINE Ad26.COV2.S- Brain haemorrhaging, Blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BRAIN HEMORRHAGE; BLOOD CLOT

VAERS ID: 1577475 (history)

Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-17

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: The patient was very healthy.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210818781

Write-up: DIED FROM THE VACCINE; This spontaneous report received from a consumer via a company representative concerned a male of unspecified age. The patient''s height, and weight were not reported. The patient was very healthy. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, after vaccination patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0-20210818781-COVID-19 Vaccine AD26.COV2.S-Died From The Vaccine. This event(s) is considered un-assessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1594327 (history)
Form: Version 2.0

Age:
Sex: Unknown
Location: Wisconsin
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210829146

Write-up: This spontaneous report received from a consumer concerned a patient of an unspecified age, sex, race and ethnic origin. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown expiry: unknown) with frequency 1 total administered, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. According to the hearsay on an unspecified date, one patient died from unknown cause of death. It was unknown if autopsy was performed. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210829146-Covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1594329 (history)
Form: Version 2.0
Age: 50.0

Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Suspected COVID-19, Thrombosis, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210829189

Write-up: DEATH; BLOOD CLOTS; SUSPECTED CLINICAL VACCINATION FAILURE; COVID RELATED ISSUES; This spontaneous report received from a company representative concerned a 50year old male. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. According to the hearsay, a patient died after taking the Janssen Covid 19 shot some time ago. On an unspecified date, the patient died with Covid related issues (suspected clinical vaccination failure) and he had blood clots. On an unspecified date, the patient died from unknown cause of death. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of death on an unspecified date, and the outcome of covid related issues, blood clots and suspected clinical vaccination failure was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender"s Comments: V0: 20210829189-COVID-19 VACCINE AD26.COV2.S-death, blood clots. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210829189-COVID-19 VACCINE AD26.COV2.S-Suspected clinical vaccination failure. This event(s) is considered not related. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown

scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1594357 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Maryland
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-21

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: It was reported that, the patient was fit.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210832457

Write-up: SUSPECTED HEART ATTACK; This spontaneous report received from a company representative concerned a 60 year old male patient. The patient''s height, and weight were not reported. The patient''s pre-existing medical conditions included: It was reported that, the patient was fit. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknow) with frequency as 1 total for prophylactic vaccination. Dose and therapy start date were not reported. The batch number was not reported. Per Procedure, no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient was suspected with heart attack. On an unspecified date, the patient died from suspected heart attack. It was reported that, the patient died within 30 days of receiving vaccine and cause of death was unknown however suspected as heart attack. It was unknown if the autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0: 20210832457- covid-19 vaccine ad26.cov2.s-suspected heart

attack. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: SUSPECTED HEART ATTACK

VAERS ID: 1594405 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-21

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Headache, Palpitations
SMQs:, Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210835168

Write-up: DIED; HEART RACING; HEADACHE; This spontaneous report received from a consumer via a company representative via social media concerned a female of unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported. Per procedure, no follow-up will be requested for this case. On an unspecified date, the patient complained of heart racing and headache and the patient died two weeks later. The cause of death was unknown. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of unknown cause of death on an unspecified date, and the

outcome of heart racing and headache was not reported. This report was serious (Death).; Sender"s Comments: V0 20210835168-COVID-19 VACCINE AD26.COV2.S-died. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1597748 (history)
Form: Version 2.0
Age: 39.0
Sex: Female
Location: Utah
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-21

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Pain in extremity
SMQs:, Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (No reported medical history)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20210

Write-up: Sore arm; This spontaneous case was reported by a consumer and describes the occurrence of PAIN IN EXTREMITY (Sore arm) in a 39-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient"s past medical history included No adverse event (No reported medical history). On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced PAIN IN EXTREMITY (Sore arm). At the time of the report, PAIN IN EXTREMITY (Sore arm) outcome was unknown. An autopsy was performed. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. This case was linked to MOD-2021-043354 (Patient

Link). Reporter did not allow further contact

VAERS ID: 1629288 (history)
Form: Version 2.0
Age: 43.0
Sex: Female
Location: California
Vaccinated: 2021-04-21
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-24

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

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Death, Malaise, SARS-CoV-2 test positive
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-21
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:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type:

Write-up: THE PATIENT CAME ON 8/16 /2021 WITH COVID SYMPTOMES TESTED WITH PCR THE RESULT CAME POSITIVE FOR COVID THE PATIENT EXPIRED ON 8/21/2021

VAERS ID: 1631068 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-08-25

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210843032

Write-up: DEATH; This spontaneous report received from a consumer via social media and concerned two male patients of unspecified age. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, 1 total administered, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patients died from unknown cause. It was unknown whether an autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patients died (due to unknown cause) on an unspecified date. This report was serious (Death).; Sender"s Comments: V0: 20210843032- Covid-19 vaccine ad26.cov2.s- Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1631079 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-08-25

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

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210844140

Write-up: DIED OF HEART ATTACK; This spontaneous report received from a consumer via a company representative via social media concerned a patient of unspecified age and sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown)1 total frequency, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. As per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient died of heart attack. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender''s Comments: V0 20210844140-COVID-19 VACCINE AD26.COV2.S-died of heart attack. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: HEART ATTACK

VAERS ID: 1631954 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Idaho
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-25

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

Administered by: Other **Purchased by:** ?
Symptoms: Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210842457

Write-up: DIED FROM COMPLICATIONS OF COVID/COVID AFTER VACCINATED; SUSPECTED CLINICAL VACCINATION FAILURE; This spontaneous report received from a physician via a company representative via social media concerned a 70 year old male. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unk) dose was not reported, 1 total, administered on MAR-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient died from complications of covid/covid after vaccinated (suspected clinical vaccination failure). Patient was fully vaccinated with the J&J vaccine. It was reported that there was lack of guidance about boosting that vaccine led to his death. It was reported that patient was pretty healthy. Cause of death was reported as complications of covid. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of died from complications of covid/covid after vaccinated on an unspecified date, and the outcome of suspected clinical vaccination failure was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender''s Comments: V0: 20210842457-Covid-19 vaccine ad26.cov2.s-died from complications of covid/covid after vaccinated.This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210842457-Covid-19 vaccine ad26.cov2. s- Suspected clinical vaccination failure. This event(s) is considered not related. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS; Reported Cause(s) of Death: DIED FROM COMPLICATIONS OF COVID

VAERS ID: 1634859 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-26

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210845181

Write-up: DEATH; This spontaneous report received from a consumer via a company representative from social media concerned an 8 decade old of unspecified age, sex, race and ethnicity. The patient"s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported, expiry: Unknown) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210844359.; Sender"s Comments: V0: 20210845181-covid-19 vaccine ad26.cov2.s-Death.This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1642962 (history)

Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Hospitalisation
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: USJNJFOC20210846858

Write-up: ENDED UP IN THE HOSPITAL; DEATH; This spontaneous report received from a consumer via a company representative from social media concerned "multiple patients". The patient"s height, and weight were not reported. No past medical histories or concurrent conditions were reported. The patients received covid-19 vaccine (suspension for injection, route of admin not reported, batch number: Unknown) dose were not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch numbers were not reported and have been requested. No concomitant medications were reported. On an unspecified date, the many patients got vaccinated and ended up in the hospital, and were hospitalized (date unspecified). The number of days of hospitalization were not reported. On an unspecified date, the patients died from an unknown cause of death. It was unspecified if an autopsy was performed. The cause of death was not reported. The action taken with covid-19 vaccine was not applicable. The patients died of death and ended up in the hospital on an unspecified date. This report was serious (Death, and Hospitalization Caused / Prolonged). This case, from the same reporter is linked to 20210844140, 20210844317 and 20210844359.; Sender"s Comments: V0-20210846858-COVID-19 VACCINE AD26.COV2.S-Death and Ended up in the Hospital(Fatal); These events are considered un-assessable. The events have a compatible/suggestive temporal relationship, are unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with

the events.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1644373 (history)
Form: Version 2.0
Age: 85.0
Sex: Female
Location: Arizona
Vaccinated: 2021-01-28
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	041L20A / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Hepatic failure, Portal vein thrombosis
SMQs:, Hepatic failure, fibrosis and cirrhosis and other liver damage-related conditions (narrow), Embolic and thrombotic events, venous (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: Diabetes; Hypertension
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: liver failure; developed a clot in her portal vein; This spontaneous case was reported by a consumer and describes the occurrence of PORTAL VEIN THROMBOSIS (developed a clot in her portal vein) and HEPATIC FAILURE (liver failure) in an 85-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 002A21A and 041L20A) for COVID-19 vaccination. Concurrent medical conditions included Hypertension and Diabetes. On 28-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 26-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced PORTAL VEIN THROMBOSIS (developed a clot in her portal vein) (seriousness criteria death, hospitalization and medically significant) and HEPATIC FAILURE (liver failure) (seriousness criteria death, hospitalization and medically significant). The reported cause of death was Liver failure. It is

unknown if an autopsy was performed. It was reported that the reporter believed that her mother passed away as a result of a complication with the vaccine. Patient''s mother received first dose on 28-JAN-2021. Patient received second dose one 26-FEB-2021. The patient was 85-year-old with hypertension and diabetes and developed a clot in her portal vein. Patient was hospitalized. The clot made her go into liver failure which was the cause of death. The date of the reported events remained unknown at the time of the report. Treatment information were not provided by the reporter. Concomitant medications were not reported by reporter. Company Comment: This is the case of an 85-year-old female subject with a history of hypetension and diabetes who died due to hepatic falure. Reportedly, the patient initially developed thrombosis in portal vein (a clot in her portal vein), however a temporal association between the use of the product and the start date of the event was not reported. The patient subsequently developed liver failure and died due to liver failure, as reported, however, the date of death was not provided. It is unknown if an autopsy was performed. Very limited information regarding these events has been provided at this time. Causality is confounded with the patient''s advanced age, hypertension and diabetes.; Sender''s Comments: This is the case of an 85-year-old female subject with a history of hypetension and diabetes who died due to hepatic falure. Reportedly, the patient initially developed thrombosis in portal vein (a clot in her portal vein), however a temporal association between the use of the product and the start date of the event was not reported. The patient subsequently developed liver failure and died due to liver failure, as reported, however, the date of death was not provided. It is unknown if an autopsy was performed. Very limited information regarding these events has been provided at this time. Causality is confounded with the patient''s advanced age, hypertension and diabetes.; Reported Cause(s) of Death: Liver failure

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VAERS ID: 1644390 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: USMODERNATX, INC.MOD20212

Write-up: Blood clots; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criteria death, hospitalization and medically significant). The reported cause of death was blood clots. It is unknown if an autopsy was performed. The concomitant medication was not reported. A consumer called and stated that his uncle passed away 2 days after receiving Moderna Covis-19 Vaccine. Reporter stated that his uncle had to go to ICU due to blood clots and died from it. No treatment medication information was provided. This is a case of death of a Male subject who passed away two days after receiving the vaccine. Very limited information regarding the event has been provided at this time. No further information is expected.; Sender''s Comments: This is a case of death of a Male subject who passed away two days after receiving the vaccine. Very limited information regarding the event has been provided at this time. No further information is expected.; Reported Cause(s) of Death: blood clots

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VAERS ID: 1644391 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Utah
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 2	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: patient died after the second dose of the Moderna vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (patient died after the second dose of the Moderna vaccine) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Concomitant product use was not provided by the reporter. Treatment information was not provided. Company Comment: Very limited information regarding the events has been provided at this time.; Sender"s Comments: Very limited information regarding the events has been provided at this time.; Reported Cause(s) of Death: patient died after the second dose of the Moderna vaccine

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VAERS ID: 1644393 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:

Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Death after vaccination; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Death after vaccination) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant medication details was provided. No treatment medication details was provided. Company comment: Very limited information regarding the events has been provided at this time. Further information is not expected.; Sender"s Comments: Very limited information regarding the events has been provided at this time. Further information is not expected.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1644452 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Death; This spontaneous case was reported by a patient family member or friend and describes the occurrence of DEATH (Death) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant products were not provided. Treatment medication were not reported. Company Comment: Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender''s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1644831 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: cardiac arrest; This spontaneous case was reported by a consumer and describes the occurrence of CARDIAC ARREST (cardiac arrest) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced CARDIAC ARREST (cardiac arrest) (seriousness criteria death and medically significant). The reported cause of death was Cardiac arrest. It is unknown if an autopsy was performed. No concomitant medications were provided by the reporter. No treatment information was provided by the reporter. This is a fatal case of cardiac arrest in a female, age unknown, who received mRNA-1273 on an unknown date. Cause of death was cardiac arrest. Very limited information has been provided at this time. Further information has been requested.; Sender"s Comments: This is a fatal case of cardiac arrest in a female, age unknown, who received mRNA-1273 on an unknown date. Cause of death was cardiac arrest. Very limited information has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: cardiac arrest

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VAERS ID: 1645002 (history)
Form: Version 2.0
Age: 65.0
Sex: Male
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Someone passed away after getting the vaccine; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Someone passed away after getting the vaccine) in a 65-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product use was not provided by the reporter. Treatment information was not provided. The patient died the day after receiving the Moderna Covid-19 vaccine. Very limited information regarding this event has been provided at this time. Further information can"t be requested (No consent for safety follow-up).; Sender"s Comments: Very limited information regarding this event has been provided at this time. Further information can"t be requested (No consent for safety follow-up).; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1645033 (history)
Form: Version 2.0
Age: 41.0
Sex: Male
Location: West Virginia
Vaccinated: 2021-06-22
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	017C21A / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Adverse event, Headache
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-07-30
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: few other things; headache; This spontaneous case was reported by an other health care professional and describes the occurrence of ADVERSE EVENT (few other things) in a 41-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine)

(batch no. 017C21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 22-Jun-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced ADVERSE EVENT (few other things) (seriousness criterion death) and HEADACHE (headache). On 30-Jul-2021, HEADACHE (headache) had resolved. The patient died on 30-Jul-2021. The cause of death was not reported. It is unknown if an autopsy was performed. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. No relevant concomitant and treatment medications were reported Very limited information regarding the events has been provided at this time. This case was linked to MOD-2021-286501 (Patient Link).; Sender"s Comments: Very limited information regarding the events has been provided at this time.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1645093 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: No adverse event (no medical history per source document).
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: People have died from the Moderna vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and

describes the occurrence of DEATH (People have died from the Moderna vaccine) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient"s past medical history included No adverse event (no medical history per source document). On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. No concomitant products were reported. No treatment information provided. Company comment: This is a case of sudden death in a patient with unknown age and gender and with unknown past medical history who died in unknown days after receiving a dose of the vaccine. Very limited information has been provided at this time.; Sender"s Comments: This is a case of sudden death in a patient with unknown age and gender and with unknown past medical history who died in unknown days after receiving a dose of the vaccine. Very limited information has been provided at this time.; Reported Cause(s) of Death: People have died from the Moderna vaccine

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VAERS ID: 1645248 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: People that died; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of

DEATH (People that died) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. The concomitant medications were not reported. The treatment information was not provided. Company Comment : This case cannot be evaluated, it seems invalid. More info has been required to assess its validity.; Sender"s Comments: This case cannot be evaluated, it seems invalid. More info has been required to assess its validity.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1645289 (history)
Form: Version 2.0
Age: 76.0
Sex: Male
Location: Iowa
Vaccinated: 2021-01-01
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Blood bicarbonate, Blood creatinine, Blood gases, Blood glucose, Brain natriuretic peptide, COVID-19, Carbon dioxide, Chest X-ray, Cough, Fibrin D dimer, Glycosylated haemoglobin, Lymphocyte count, Oxygen saturation, PO2, Pyrexia, Respiratory failure, Respiratory rate, SARS-CoV-2 antibody test, SARS-CoV-2 test, Tachypnoea, Troponin, Vital signs measurement, White blood cell count, pH body fluid
SMQs:, Anaphylactic reaction (broad), Asthma/bronchospasm (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), Eosinophilic pneumonia (broad), Hypersensitivity (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? Yes
  Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 7 days
  Extended hospital stay? No

Previous Vaccinations:
Other Medications: TACROLIMUS; MYCOPHENOLATE MOFETIL
Current Illness: End stage renal disease; Exertional dyspnea; IgA nephropathy; Immunosuppression (on stable maintenance immunosuppression); Noninvasive mechanical ventilation (Shortly after admission, noninvasive positive pressure ventilation was initiated for increasing oxygen requirements).
Preexisting Conditions: Medical History/Concurrent Conditions: Renal transplant (with antithymocyte globulin induction 6 years prior presented with exertional dyspnea, posttransplant course has been uncomplicated without kidney allograft rejection, chronic infection, or hospitalization).
Allergies:
Diagnostic Lab Data: Test Date: 2021; Test Name: HCO3; Result Unstructured Data: 24.5 mmol/L on presenting to the hospital second time; Test Date: 2021; Test Name: serum creatinine; Result Unstructured Data: baseline serum creatinine ranging from 0.9 to 1.0 mg/dL without proteinuria.; Test Date: 2021; Test Name: serum creatinine; Result Unstructured Data: serum creatinine, 1.5 mg/dL on admission; Test Date: 2021; Test Name: arterial blood gases; Result Unstructured Data: Arterial blood gas (FiO2 0.44) confirmed acute hypoxic respiratory failure on presenting to the hospital second time; Test Date: 2021; Test Name: Non-fasting glucose; Result Unstructured Data: Non-fasting glucose was 492 mg/dL; Test Date: 2021; Test Name: BNP; Result Unstructured Data: BNP, 175 pg/mL on admission; Test Date: 2021; Test Name: PaCO2; Result Unstructured Data: 30.3 mm Hg on presenting to the hospital second time; Test Date: 2021; Test Name: Chest X-ray; Result Unstructured Data: Chest X-ray did not demonstrate any acute findings; Test Date: 2021; Test Name: Chest X-ray; Result Unstructured Data: Chest X-ray now showed an interstitial opacity in the left lower lung on presenting to the hospital second time; Test Date: 2021; Test Name: D-dimer; Result Unstructured Data: D-dimer, 557 ng/mL on admission; Test Date: 2021; Test Name: hemoglobin A1c; Result Unstructured Data: hemoglobin A1c was 8.2%; Test Date: 2021; Test Name: absolute lymphocyte count; Result Unstructured Data: 150/?L (baseline 5 months prior: 550/?L) - On admission; Test Date: 2021; Test Name: pulse oximeter; Result Unstructured Data: pulse oximeter reading was 66% on ambient air on presenting to the hospital second time; Test Date: 2021; Test Name: SatO2; Result Unstructured Data: 89.7% despite supplemental oxygen on presenting to the hospital second time; Test Date: 2021; Test Name: pH; Result Unstructured Data: pH 7.49 on presenting to the hospital second time; Test Date: 2021; Test Name: PaO2; Result Unstructured Data: 55.5mm Hg on presenting to the hospital second time; Test Date: 2021; Test Name: respiratory rate; Result Unstructured Data: 30 bpm on presenting to the hospital second time; Test Date: 2021; Test Name: SARS-CoV-2 antispike antibody; Test Result: Negative; Result Unstructured Data: A SARS-CoV-2 antispike antibody was undetectable (<0.8 units) by semiquantitative immunoassay; Comments: used Elecsys Anti-SARS-CoV-2 S, ; Test Date: 2021; Test Name: SARS-CoV-2; Test Result: Positive; Result Unstructured Data: COVID-19 was confirmed with a positive nasopharyngeal swab sample by SARS-CoV-2 real-time reverse transcription-polymerase chain reaction with a cycle threshold value for target N2 of 19.5; Comments: Used Xpert Xpress SARS-CoV-2, Cepheid; Test Date: 2021; Test Name: SARS-CoV-2; Test Result: Positive; Result Unstructured Data: A repeat nasopharyngeal sample obtained 2 days after admission confirmed persistent positivity of SARS CoV- 2 by reverse transcription-polymerase chain reaction with a cycle threshold value for target N2 gene of 15.2; Test Date: 2021; Test Name: Troponin; Result Unstructured Data: Troponin-I, 0.01 ng/mL on admission; Test Date: 2021; Test Name: Vital signs; Result Unstructured Data: Vital signs were notable for tachypnea: body temperature 36.7 ?C, blood pressure 127/83 mm Hg, heart rate 83 bpm, respiratory rate 24 bpm, and SatO2 96% on ambient air.; Comments: Patient"s physical examination was otherwise unremarkable; Test Date: 2021; Test Name: White blood cell count; Result Unstructured Data: White blood cell count was 7500/?L on admission.
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: fatal COVID-19 infection; hypoxic respiratory failure; tachypnea/tachypneic; nonproductive cough; intermittent low-grade; This literature-non-study case was reported in a literature article and describes the occurrence of COVID-19 (fatal COVID-19 infection), RESPIRATORY FAILURE (hypoxic respiratory failure) and TACHYPNOEA (tachypnea/tachypneic) in a 76-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. LITERATURE REFERENCE: Case report: Severe COVID-19 in a kidney transplant recipient without humoral response to SARS-CoV-2 mRNA vaccine series. Transplant Direct. 2021;7(9):e473 The patient''s past medical history included Renal transplant (with antithymocyte globulin induction 6 years prior presented with exertional dyspnea, posttransplant course has been uncomplicated without kidney allograft rejection, chronic infection, or hospitalization.) in 2015 and Noninvasive mechanical ventilation (Shortly after admission, noninvasive positive pressure ventilation was initiated for increasing oxygen requirements.) since an unknown date. Concurrent medical conditions included End stage renal disease, IgA nephropathy, Immunosuppression (on stable maintenance immunosuppression) and Exertional dyspnea. Concomitant products included TACROLIMUS and MYCOPHENOLATE MOFETIL for Immunosuppression. In January 2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. In February 2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced COVID-19 (fatal COVID-19 infection) (seriousness criteria death and hospitalization prolonged), RESPIRATORY FAILURE (hypoxic respiratory failure) (seriousness criteria death, hospitalization prolonged and medically significant), TACHYPNOEA (tachypnea/tachypneic) (seriousness criterion hospitalization prolonged), COUGH (nonproductive cough) and PYREXIA (intermittent low-grade). The patient was hospitalized for 7 days due to COVID-19, RESPIRATORY FAILURE and TACHYPNOEA. The patient was treated with DEXAMETHASONE (intravenous) for COVID-19, at a dose of 6 mg once a day; REMDESIVIR (intravenous) at a dose of 100 mg once a day; PREDNISONE for Cough and Fever, at a dose of 30 milligram; PREDNISONE for Cough and Fever, at a dose of 20 milligram and PREDNISONE for Cough and Fever, at a dose of 10 milligram. The patient died on an unknown date. The reported cause of death was this case of a kidney transplant recipient who developed a fatal covid-19 infection about 11 d after completion of the sars-cov-2 mrna vaccine series and Hypoxic respiratory failure. It is unknown if an autopsy was performed. At the time of death, TACHYPNOEA (tachypnea/tachypneic), COUGH (nonproductive cough) and PYREXIA (intermittent low-grade) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In 2021, Blood bicarbonate: high (High) 24.5 mmol/L on presenting to the hospital second time. In 2021, Blood creatinine: 0.9 to 1.0 mg/dl (normal) baseline serum creatinine ranging from 0.9 to 1.0 mg/dL without proteinuria. and normal (normal) serum creatinine, 1.5 mg/dL on admission. In 2021, Blood gases: abnormal (abnormal) Arterial blood gas (FiO2 0.44) confirmed acute hypoxic respiratory failure on presenting to the hospital second time. In 2021, Blood glucose: high (High) Non-fasting glucose was 492 mg/dL. In 2021, Brain natriuretic peptide: normal (normal) BNP, 175 pg/mL on admission. In 2021, Carbon dioxide: low (Low) 30.3 mm Hg on presenting to the hospital second time. In 2021, Chest X-ray: normal (normal) Chest X-ray did not demonstrate any acute findings and abnormal (abnormal) Chest X-ray now showed an interstitial opacity in the left lower lung on presenting to the hospital second time. In 2021, Fibrin D dimer: 557 ng/ml (abnormal) D-dimer, 557 ng/mL on admission. In 2021, Glycosylated haemoglobin: high (High) hemoglobin A1c was 8.2%. In 2021, Lymphocyte count: low (Low) 150/?L (baseline 5 months prior: 550/?L) - On admission. In 2021, Oxygen saturation: low (Low) pulse oximeter reading was 66% on ambient air on presenting to the hospital second time and low (Low) 89.7% despite supplemental oxygen on presenting to the hospital second time. In 2021, PO2: low (Low) 55.5mm Hg on presenting to the hospital second time. In 2021, Respiratory rate: high (High) 30 bpm on presenting to the hospital second time. In 2021, SARS-CoV-2 antibody test: negative (Negative) A SARS-CoV-

2 antispike antibody was undetectable (<0.8 units) by semiquantitative immunoassay. In 2021, SARS-CoV-2 test: positive (Positive) COVID-19 was confirmed with a positive nasopharyngeal swab sample by SARS-CoV-2 real-time reverse transcription-polymerase chain reaction with a cycle threshold value for target N2 of 19.5 and positive (Positive) A repeat nasopharyngeal sample obtained 2 days after admission confirmed persistent positivity of SARS CoV- 2 by reverse transcription-polymerase chain reaction with a cycle threshold value for target N2 gene of 15.2. In 2021, Troponin: low (Low) Troponin-I, 0.01 ng/mL on admission. In 2021, Vital signs measurement: normal (normal) Vital signs were notable for tachypnea: body temperature 36.7 ?C, blood pressure 127/83 mm Hg, heart rate 83 bpm, respiratory rate 24 bpm, and SatO2 96% on ambient air.. In 2021, White blood cell count: normal (normal) White blood cell count was 7500/?L on admission. In 2021, pH body fluid: high (High) pH 7.49 on presenting to the hospital second time. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter considered COVID-19 (fatal COVID-19 infection), RESPIRATORY FAILURE (hypoxic respiratory failure), TACHYPNOEA (tachypnea/tachypneic), COUGH (nonproductive cough) and PYREXIA (intermittent low-grade) to be possibly related. Patient started to develop nonproductive cough and intermittent low-grade fevers. His symptoms gradually worsened, and 5 days before admission, he was evaluated in the emergency department. Patient was discharged with a 6 day course of glucocorticoid therapy. He was not tested for SARS-CoV-2 at that time because he had been fully vaccinated. Patient was admitted to the intensive care unit for further treatment after presenting to the hospital the second time where he was found to be tachypneic and severely hypoxic. Upon admission, his exam was notable for bibasilar crackles in lower lung fields without peripheral edema. Patient continued to decompensate and died on the seventh day after admission. Company Comment: Very limited information regarding this event/s has been provided at this time. Further information has been requested. Further information is not expected.; Sender"s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested. Further information is not expected.; Reported Cause(s) of Death: This case of a kidney transplant recipient who developed a fatal COVID-19 infection about 11 d after completion of the SARS-CoV-2 mRNA vaccine series; hypoxic respiratory failure

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VAERS ID: 1645361 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Heart rate increased
SMQs:, Neuroleptic malignant syndrome (broad), Arrhythmia related investigations, signs and symptoms (broad), Dehydration (broad)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: died from a rapid heartbeat after getting the vaccine; Result Unstructured Data: increased
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: died from a rapid heartbeat after getting the vaccine; This spontaneous case was reported by a consumer and describes the occurrence of HEART RATE INCREASED (died from a rapid heartbeat after getting the vaccine) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced HEART RATE INCREASED (died from a rapid heartbeat after getting the vaccine) (seriousness criterion death). The reported cause of death was died from a rapid heartbeat after getting the vaccine. It is unknown if an autopsy was performed. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, Heart rate increased: increased (High) increased. No concomitant medication details was reported. No treatment medication details was reported. Company Comment: This case concerns a female patient of unknown age who had a fatal event of HEART RATE INCREASED after receiving mRNA-1273. The reported cause of death was died from a rapid heartbeat after getting the vaccine. It is unknown if an autopsy was performed. Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender"s Comments: This case concerns a female patient of unknown age who had a fatal event of HEART RATE INCREASED after receiving mRNA-1273. The reported cause of death was died from a rapid heartbeat after getting the vaccine. It is unknown if an autopsy was performed. Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: died from a rapid heartbeat after getting the vaccine

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VAERS ID: 1645370 (history)
Form: Version 2.0
Age: 72.0
Sex: Male
Location: Mississippi
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 3	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Fall
SMQs:, Accidents and injuries (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-20
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Fell down on the floor and died; This spontaneous case was reported by a nurse and describes the occurrence of FALL (Fell down on the floor and died) in a 72-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received third dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced FALL (Fell down on the floor and died) (seriousness criterion death). The patient died on 20-Aug-2021. The reported cause of death was fell on the floor and died. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant product use was not provided by the reporter. The patient died yesterday after 9 days [unsure of the date] of getting the booster dose of the Moderna COVID-19 vaccine. The patient did not had any side effects with the first 2 doses of the Moderna COVID-19 Vaccine. Treatment information was not provided. Company Comment : Very limited information regarding the events has been provided at this time. Further information has been requested.; Sender"s Comments: Very limited information regarding the events has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Fell on the floor and died

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VAERS ID: 1645387 (history)
Form: Version 2.0
Age: 42.0
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	- / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Angiocardiogram, Angiogram, Autopsy, Cardiogenic shock, Chest pain, Dyspnoea, Echocardiogram, Electrocardiogram, Myocarditis, Pyrexia, SARS-CoV-2 test, Serology test, Tachycardia, Ventricular dysfunction
SMQs:, Cardiac failure (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) (narrow), Acute central respiratory depression (broad), Pulmonary hypertension (broad), Gastrointestinal nonspecific symptoms and therapeutic procedures (broad), Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Dehydration (broad), Immune-mediated/autoimmune disorders (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Date: 2021; Test Name: Coronary angiography; Result Unstructured Data: revealed no coronary artery disease.; Test Date: 2021; Test Name: Coronary angiography; Result Unstructured Data: revealed no coronary artery disease.; Test Date: 2021; Test Name: Autopsy; Result Unstructured Data: An inflammatory infiltrate admixed with macrophages, T-cells, eosinophils, and B cells was observed; Test Date: 2021; Test Name: transthoracic Echocardiogram; Result Unstructured Data: global biventricular dysfunction (ejection fraction, 15%), normal ventricular dimensions, and left ventricular hypertrophy; Test Date: 2021; Test Name: Electrocardiogram; Result Unstructured Data: showed diffuse ST-segment elevation; Test Date: 2021; Test Name: SARS-CoV-2 PCR test; Test Result: Negative ; Result Unstructured Data: Negative for SARS-CoV-2 and no other causes were identified.; Test Date: 2021; Test Name: serologic examination; Result Unstructured Data: no other causes were identified.
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Cardiogenic shock; biventricular myocarditis; Global biventricular dysfunction; tachycardia; chest pain; dyspnea; Fever; This literature-non-study case was reported in a literature article and describes the occurrence of CARDIOGENIC SHOCK (Cardiogenic shock), MYOCARDITIS (biventricular myocarditis) and VENTRICULAR DYSFUNCTION (Global biventricular dysfunction) in a 42-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced CARDIOGENIC SHOCK (Cardiogenic shock)

(seriousness criteria death and medically significant), MYOCARDITIS (biventricular myocarditis) (seriousness criteria death and medically significant), VENTRICULAR DYSFUNCTION (Global biventricular dysfunction) (seriousness criterion death), TACHYCARDIA (tachycardia), CHEST PAIN (chest pain), DYSPNOEA (dyspnea) and PYREXIA (Fever). The patient died on an unknown date. The reported cause of death was biventricular myocarditis, Cardiogenic shock and global biventricular dysfunction. An autopsy was performed. At the time of death, TACHYCARDIA (tachycardia), CHEST PAIN (chest pain), DYSPNOEA (dyspnea) and PYREXIA (Fever) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In 2021, Angiocardiogram: normal (normal) revealed no coronary artery disease.. In 2021, Angiogram: normal (normal) revealed no coronary artery disease.. In 2021, Autopsy: biventricular myocarditis (abnormal) An inflammatory infiltrate admixed with macrophages, T-cells, eosinophils, and B cells was observed. In 2021, Echocardiogram: abnormal (abnormal) global biventricular dysfunction (ejection fraction, 15%), normal ventricular dimensions, and left ventricular hypertrophy. In 2021, Electrocardiogram: abnormal (abnormal) showed diffuse ST-segment elevation. In 2021, SARS-CoV-2 test: negative (Negative) Negative for SARS-CoV-2 and no other causes were identified.. In 2021, Serology test: abnormal (abnormal) no other causes were identified.. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Patient had not reported a viral prodrome The letter stated that a direct causal relationship could not be definitively established because they did not perform testing for viral genomes or autoantibodies in the tissue specimens. However, no other causes were identified by PCR assay or serologic examination. Concomitant product use was not provided by the reporter. ? No treatment information was provided. Very limited information regarding this event/s has been provided at this time. Further information has been requested. Further information is not expected.; Sender''s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested. Further information is not expected.; Reported Cause(s) of Death: biventricular myocarditis; Cardiogenic shock; Global biventricular dysfunction

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VAERS ID: 1645838 (history)
Form: Version 2.0
Age: 52.0
Sex: Female
Location: Arizona
Vaccinated: 2021-05-31
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Pharmacy **Purchased by:** ?
Symptoms: Crying
SMQs:, Depression (excl suicide and self injury) (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Asthma; Death (Patient mother); Death (Patient father); Emotional suffering (stroke 20 years ago and it effected her emotions. could only laugh or be mad.); Heart attack (This was eight years ago.); Kidney disorder; Lung disorder; Schizophrenia; Stent placement (capillaries were small.The veins to he heart were small.); Stroke (Occurred 20 years ago)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100965474

Write-up: After she first got the shot she started crying.; This is a spontaneous report from a contactable consumer or other non hcp (patient''s husband). A 52-years-old female patient received first dose of bnt162b2 (BNT162B2, PFIZER-BIOECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration, administered in Arm Left on 31May2021 (Age at vaccination: 52-years-old) as single dose for covid-19 immunization. (Vaccination facility: Pharmacy/Drug Store). Medical history included stroke occurred 20 years ago, kidney problems, heart attack eight years ago, patient had 7 stents around her heart, her capillaries were small, The veins to her heart were small, asthma, schizophrenia, stated lungs were bad, had a stroke 20 years ago and it affected her emotions. She could only laugh or be mad. family medical history included Patient mother and father had passed away within three months of each other. The patient concomitant medications were not reported. patient did not receive any other vaccine within 4 weeks Prior Vaccinations. Caller on the line who mentioned he was a PCA, clarified as a patient care attendant for his wife. He is calling about the Pfizer Shot, the vaccination, clarified as to prevent COVID, the COVID Shot. He stated his wife died due to the shot. He mentioned he has no avenue to go down. No direction. He stated he is upset no research has been done form people who died from the shot, or information from it causing one to have a heart attack, lungs filling with water, or weakening the immune system. He later clarified this is what happened to his wife. Caller stated he feels that his wife died from the COVID shot, but he is not a scientist and has no way of proving that. He wants someone to call him and tell him. Caller clarified the patient''s cause of death. He stated the death certificate has she passed away due to heart attack due to lungs filling with water and she could not be resuscitated. They tried for two hours. Caller stated he does not know the exact date patient passed away, it is all packed up. It was either 02Jul2021 or 03Jul2021. He confirmed the heart attack and lungs filling with water occurred on the date she passed away. Caller explained, he wanted to explain what he saw. After she first got the shot she started crying. He mentioned his wife had a stroke 20 years ago and it affected her emotions. She could only laugh or be mad. Patient''s mother and father had passed away within three months of each other and she could not even cry then. Caller stated after patient got the shot she started crying and did not know what was going on. He mentioned she was close to menopause and he thought it was just menopause going on. It was all day, every day, she was crying. Then after the second shot a few days later patient stopped breathing. The paramedics were called who provided a breathing treatment. Patient was taken to the hospital and was there for three days. They could not figure out what was wrong. They did work-ups. Patient was in ICU. Then she crashed. They started CPR. Patient''s lungs filled up with water, patient had a heart attack, and she stopped breathing. Caller clarified patient had the issues with breathing and stopped breathing initially

like on 28Jun2021. Then patient was taken to the hospital. Stated patient''s health was never like us. However, he kept her not sick. He was good at his job. However, when patient was sick he would take patient to the hospital or the doctor''s to find out what is going on. When sick she was in the hospital, when she was fine she was home. No vaccines administered on same date of the pfizer suspect. Confirmed none were given on the same date. The paramedics were called, patient was taken to hospital and was admitted to ICU due to patient stopped breathing. The outcome for the event was reported as unknown at the time of his report. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.

VAERS ID: 1646195 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 2021-05-03
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	ER8735 / 2	- / OT

Administered by: Private **Purchased by:** ?
Symptoms: Maternal exposure during pregnancy, Premature baby
SMQs:, Pregnancy, labour and delivery complications and risk factors (excl abortions and stillbirth) (narrow), Neonatal disorders (narrow)

Life Threatening? Yes
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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 INC202100987373

Write-up: Preemie baby died in the NICU; Maternal Exposure During Pregnancy; This is a spontaneous report from a contactable consumer (patient''s mother). This consumer reported information for both mother and fetus/baby. This is a baby report. A 36-year-old female

pregnant patient received BNT162B2, first dose via an unspecified route of administration, administered in right arm on 12Apr2021 05:00 PM (Batch/Lot Number: EW0153) as dose 1, single; second dose via an unspecified route of administration, administered in left arm on 03May2021 (Batch/Lot Number: ER8735) as dose 2, single for covid-19 immunization. The patient"s medical history was not reported. There were no concomitant medications. Until Monday the 17, She has emergency cesarean to deliver at 24 weeks, 4 weeks later the preemie baby died in the NICU. The preemie baby died in the NICU in 2021. The event was serious per death, life threatening, hospitalization. The patient died in 2021. It was not reported if an autopsy was performed.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202100986436 mother case;US-PFIZER INC-202100987372 mother case; Reported Cause(s) of Death: Maternal exposure during pregnancy; 4 weeks later our preemie baby died in the NICU

VAERS ID: 1646213 (history)
Form: Version 2.0
Age: 88.0
Sex: Female
Location: North Carolina
Vaccinated: 2021-03-25
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Confusional state, Herpes zoster, Inappropriate schedule of product administration, Vaccination complication
SMQs:, Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Dementia (broad), Noninfectious encephalitis (broad), Noninfectious encephalopathy/delirium (broad), Noninfectious meningitis (broad), Medication errors (narrow), Hypoglycaemia (broad), Opportunistic infections (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-05-07
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: JAKAFI
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Myeloid leukemia
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100991285

Write-up: complete state of confusion and disillusion; Dose 1 on 05Feb2021, doe 2 on 25Mar2021; shingles; acute reaction to the vaccine and it killed her/ the shot killed her; This is a spontaneous report from a contactable consumer. An 88-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on 25Mar2021 (at the age of 88 years old) (Batch/Lot number was not reported) as single dose for covid-19 immunisation. Medical history included Myeloid leukaemia from early 2020. Concomitant medication included ruxolitinib phosphate (JAKAFI) taken for myeloid leukaemia. The patient previously received the first dose of BNT162B2 on 05Feb2021 (at the age of 88 years old) for Covid-19 Immunization and experienced confused state; panicked; Urinary tract infection (UTI); shingles; within 2 days it looked like somebody had punched her in her face, her face was purple on one side; shooting pain, across left cheek from her nose, her cheeks, down to her mouth. On 25Mar2021, somebody at that recovery center thought that it was a good idea for her to go get another vaccine. And the patient also had acute myeloid leukemia (Later clarified as history) and this was known by all parties. And when she got that vaccine on 25Mar2021, she didn''t even make it back to the recovery center, they took her straight to the hospital because she went in to a complete state of confusion and disillusion then. And she really never recovered and she died on 07May2021 and the reporter just want to let you all know because reporter just listened to NPR and listened to doctors try to prove some relationship between the shingles and the vaccine, and reporter was not saying shingles caused it or vice versa but she had an acute reaction to the vaccine and it killed her. For the patient''s Height and Weight, Reporter stated She was 4 foot 11, she was a little bit taller but as she got older her spinal cord begin to kind of shrink up on her, so 4 foot 11 works, she was barely 100 pounds when she died. For the Reason of Death, Reporter stated Somebody decided that it was a good idea to give her second COVID vaccine, that''s what killed her. If she hadn''t got the second COVID vaccine, she was recovering, she was doing much better, and the reporter spoke to her on 24Mar2021, just before she got that shot and she was never the same after that, the shot killed her, reporter was not saying you all killed her, she was saying the person who said it was a good idea to give her the shot, killed the patient. The reporter was not a doctor, resisted it somebody not do that but they went ahead and did it anyway and it killed her. She suffered for 90 days after she got her shot and suffered like hell after she got her second shot. It was unknown if Autopsy was performed. The outcome of event "acute reaction to the vaccine and it killed her/ the shot killed her" was fatal, of the other events was unknown. The lot number for the vaccine bnt162b2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: acute reaction to the vaccine and it killed her/ the shot killed her

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VAERS ID: 1646432 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 2021-05-01
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202100999479

Write-up: died; This is a spontaneous report from a contactable healthcare professional via medical information team and via a Pfizer-sponsored program. A 55-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration in May2021 (Batch/Lot number was not reported) as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The healthcare professional reported that they were definitely seeing a rise in COVID and that a 55-year-old guy died in their hospital the other day (unspecified date), he was vaccinated in May with Pfizer. It was unknown if an autopsy was done. Information on the lot/batch number has been requested.; Sender''s Comments: Based on the current available information and the plausible drug-event temporal association, a possible contributory role of the suspect product BNT162B2 to the development of serious event Death cannot be totally excluded. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics, concomitant medications and dates of vaccination.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.; Reported Cause(s) of Death: died

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VAERS ID: 1646449 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101000032

Write-up: developed it (blood clots) and died; This is a spontaneous report from a Pfizer-sponsored program from a non-contactable consumer or other non hcp. A female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) dose number unknown, single for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient developed it (blood clots) and died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible, information about lot/batch number cannot be obtained.; Reported Cause(s) of Death: developed it (blood clots) and died

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VAERS ID: 1646584 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Illness
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101008353

Write-up: got sick after getting vaccinated and a week later he was buried; This is a spontaneous report from a contactable consumer. This consumer (patient"s friend) reported for a male patient (reporter"s friend). A male patient of an unspecified age received bnt162b2 (BNT162B2), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. The patient got sick after getting vaccinated and a week later he was buried on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: got sick after getting vaccinated and a week later he was buried

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VAERS ID: 1646782 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-06
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101022590

Write-up: Her aunt just had the Pfizer Covid vaccine but ended up getting the delta variant then died; Her aunt just had the Pfizer Covid vaccine but ended up getting the delta variant then died; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported for a female patient (aunt) of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The medical history and concomitant medications were not reported. The consumer reported her aunt just had the PFIZER covid vaccine but ended up getting the delta variant then died. The patient died on 06Aug2021. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: Her aunt just had the Pfizer Covid vaccine but ended up getting the delta variant then died; Her aunt just had the Pfizer Covid vaccine but ended up getting the delta variant then died

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VAERS ID: 1646847 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Illinois
Vaccinated: 2021-03-01
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101024552

Write-up: developed widespread bloodclots; This is a spontaneous report from a contactable consumer for female patient (mother). A female patient of an unspecified age received bnt162b2 (COVID-19 vaccine), dose 2 via an unspecified route of administration on Mar2021 (Batch/Lot number was not reported) as dose 2, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Historical Vaccine included first dose bnt162b2 for covid-19 immunisation on Feb2021. The patient developed widespread bloodclots on 2021 with fatal outcome. The patient died on 2021. It was not reported if an autopsy was performed. The clinical course was reported as follows: patient received the COVID-19 vaccine and developed widespread blood clots. Her first dose was in February, her second dose in March and 13 days later she was gone. The lot number for the vaccine, bnt162b2, was not provided and will be requested during follow up.; Reported Cause(s) of Death: Developed widespread bloodclots

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VAERS ID: 1646859 (history)
Form: Version 2.0
Age: 61.0
Sex: Female
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route

COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Cytokine storm, Inflammation, Myocardial infarction
SMQs:, Myocardial infarction (narrow), Embolic and thrombotic events, arterial (narrow), Hypersensitivity (broad), Tumour lysis syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101026335

Write-up: heart attack; cytokine storm; inflammation; This is a spontaneous report from a contactable consumer reporting for a friend. A 61-years-old female patient received bnt162b2 (BNT162B2) at the age of 61-years-old, dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE 2, SINGLE for covid-19 immunisation. The patient medical history was not reported. The patient''s concomitant medications were not reported. Historical Vaccine included first dose of BNT162B2 for COVID-19 immunization at the age of 61 years. After the second dose, the patient had inflammation, a cytokine storm, a heart attack and then died on an unspecified date. It was not reported if an autopsy was performed. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: heart attack; cytokine storm; inflammation

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VAERS ID: 1646861 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

PFIZER/BIONTECH		

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101026586

Write-up: killed my roommate; This is a spontaneous report from a Pfizer- sponsored program. A Non-contactable consumer reported for a patient (reporters friend) that a patient of unspecified age and gender received bnt162b2 (BNT162B2, the Pfizer COVID Vaccination), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose number unknown, single for covid-19 immunisation. The patient medical history was not reported. The patient"s concomitant medications were not reported. The consumer reported that "you killed my roommate, My friends death is worth nothing to you" on an unspecified date. The patient died on an unspecified date. The outcome of the event was fatal. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: killed my roommate

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VAERS ID: 1646975 (history)
Form: Version 2.0
Age:
Sex: Male
Location: New Jersey
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?

Symptoms: Cardiac disorder, Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101033683

Write-up: passed away; heart problem; This is a spontaneous report from a contactable consumer. A 73-year-old male patient (reporter"s father) received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose 1, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient passed away from taking Covid vaccine. He never had a heart problem until he took the first Covid vaccine shot. The outcome of event heart problem was unknown. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for the vaccine, bnt162b2, was not provided and will be requested during follow up.; Reported Cause(s) of Death: passed away

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VAERS ID: 1647045 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Tennessee
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac disorder
SMQs:

Life Threatening? No

Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101036031

Write-up: heart problems; This is a spontaneous report from a contactable consumer (not the patient) via telephonic activity. A 43-year-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) (batch/lot number was not reported) via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. Historical vaccine included first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date for COVID-immunization. The patient experienced heart problems (seriousness criteria: death, medically significant) on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for the BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), was not provided and will be requested during follow up.; Reported Cause(s) of Death: heart problems

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VAERS ID: 1647060 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Michigan
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Private **Purchased by:** ?
Symptoms: Cerebrovascular accident
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)

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101038521

Write-up: female died 3 days after receiving the first COVID vaccine shot of a massive stroke; This is a spontaneous report from two contactable consumers. A 81-year-old female patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose 1, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. It was reported the female died 3 days after receiving the first COVID vaccine shot of a massive stroke. Healthy patient. It was not reported if an autopsy was performed. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.; Reported Cause(s) of Death: female died 3 days after receiving the first COVID vaccine shot of a massive stroke.

VAERS ID: 1647160 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101042649

Write-up: passed away from the Delta Variant; passed away from the Delta Variant; This is a spontaneous report from a contactable consumer (relative) from a Pfizer-sponsored program . A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) dose 1 and 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient daughter had just received an email from Pfizer stating that the patient should come in for a third dose to cover for the Delta Variant. The patient passed away from the Delta Variant on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for BNT162B2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: passed away from the Delta Variant; passed away from the Delta Variant

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VAERS ID: 1647304 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiovascular disorder, Internal haemorrhage
SMQs:, Haemorrhage terms (excl laboratory terms) (narrow), Cardiomyopathy (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-15
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No

ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101057880

Write-up: internal bleeding; cardio issues; This is a spontaneous report from a non contactable consumer. An 86-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. It was reported: "Consumer calling about her Mother in law dying from the side effects of the Pfizer Covid Vaccine at 3:00 am Sunday, 15Aug2021. Caller asked How many deaths have there been called that they believe was caused by the Pfizer vaccine? My MIL was 86 y/o and she had just started PT in the hospital. She started having cardio issues and internal bleeding and now she is dead. She had started PT and was up walking, so her doctor suggested she get the this because she was having PT, then she started having internal bleeding and cardio issues and now she"s dead. With in 48 hours of taking that vaccine. She developed symptoms where there was blood gushing out of her mouth and my father in law saw this. They couldn"t stop the bleeding because of her health condition." The patient experienced internal bleeding, cardio issues on an unspecified date. The outcome of events was fatal. The patient died on 15Aug2021. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: internal bleeding; cardio issues

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VAERS ID: 1647314 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00

Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101058096

Write-up: Died from the Pfizer Covid-19 vaccine; This is a spontaneous report from a non-contactable consumer or other non hcp. A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunization. The patient medical history was not reported. The patient''s concomitant medications were not reported. The patient died from the pfizer covid-19 vaccine on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: Died from the Pfizer Covid-19 vaccine

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VAERS ID: 1647364 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101067685

Write-up: died within 24 hours of the vaccine; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE Solution for injection), via an unspecified route of administration, on an unspecified date (Batch/lot number and expiration dates unknown), as dose number unknown, single, for COVID-19 immunisation. Relevant medical history and concomitant medications were not reported. On an unspecified date, the patient died within 24 hours of the vaccine. The patient never had a problem in his life. The reporter stated that the patient died from the vaccine. It was unknown if an autopsy was done. The lot number for the vaccine BNT162B2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: died within 24 hours of the vaccine

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VAERS ID: 1647372 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101068945

Write-up: another person died; This is a spontaneous report from a contactable consumer. A patient of unspecified gender and age received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on unspecified date at unknown dose single for COVID-19 immunization. Medical history includes historical vaccine. No concomitant medications reported. Reporter reports that after receiving the second dose, another person died. The outcome of Event was Fatal. Information about lot/batch number has been requested. Follow-up attempts have been completed, Batch/lot number not available. No further information is expected.; Reported Cause(s) of Death: another person died

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VAERS ID: 1647415 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC202101082857

Write-up: fully vaccinated father died from COVID-19 / lack of efficacy; fully vaccinated father died from COVID-19 / lack of efficacy; This is a spontaneous report from a non-contactable consumer. This consumer reported for an adult male patient received bnt162b2 (COVID-19 vaccine), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose, dose 1 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The medical history and concomitant medications were not reported. The patient experienced lack of efficacy. The reporter''s fully vaccinated father died from COVID-19. The patient died on an unspecified date. It was not reported if an autopsy was performed. Doctors said the condition of the father could have been worse if he was not vaccinated at all. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: fully vaccinated father died from COVID-19 / lack of efficacy; fully vaccinated father died from COVID-19 / lack of efficacy

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VAERS ID: 1647426 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-28

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	UNKNOWN / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC202101093096

Write-up: died two weeks ago after his 2nd dose; This is a spontaneous report from a non-contactable consumer. A male patient of unspecified age received the 2nd dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on an unspecified date as single dose for COVID-19 Immunization. Medical history and concomitant medications were unknown. The patient previously received the 1st dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) for COVID-19 Immunization. The reporter knew their personal situation where a man died two weeks ago after his 2nd dose so, the reporter was horrified that they will not considered the safety of the public before taking that emergency FDA approval because they know that it took more than just a minute of the time to determine the safety and the reporter would hope that have the integrity too consider that over the money at this point in time. When probed if the concern is with Pfizer COVID-19 vaccine, reporter stated, "Yes of course." It was unknown if autopsy was performed. Outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: died two weeks ago after his 2nd dose

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VAERS ID: 1654999 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Military **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Blood pressure abnormal; High cholesterol.
Allergies:
Diagnostic Lab Data:

CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: Friend died 2 days after receiving the vaccine; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Friend died 2 days after receiving the vaccine) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Treatment information not provided. Concomitant medications were reported as statins and blood pressure medication. This a report of dead, two days after Moderna COVID-19 vaccination in a patient of an unknown age and gender. Autopsy was not performed. Very limited information regarding the event has been provided for inferring causality. Further information is not expected; Sender''s Comments: This a report of dead, two days after Moderna COVID-19 vaccination in a patient of an unknown age and gender. Autopsy was not performed. Very limited information regarding the event has been provided for inferring causality. Further information is not expected; Reported Cause(s) of Death: unknown cause of death.

VAERS ID: 1655100 (history)
Form: Version 2.0
Age: 13.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101091051

Write-up: dead from second dose of Pfizer Covid 19 vaccine; This is a spontaneous report from a Pfizer-sponsored program by a non-contactable consumer. This report reported same event for two patients. This is the first dose of two reports. A 13-year-old female patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE 2, SINGLE at the age of 13-year-old for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient previously received the first dose of bnt162b2 (BNT162B2) for covid-19 immunisation. A 13 years old female is dead from second dose of Pfizer Covid 19 vaccine. Both had no prior conditions with the heart and now are dead. The patient died on an unspecified date. It was unknown if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101091793 same report/drug/AE, different patients; Reported Cause(s) of Death: dead from second dose of Pfizer Covid 19 vaccine

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VAERS ID: 1655104 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebral thrombosis
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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101091230

Write-up: blood clot to the brain; This is a spontaneous report from a contactable consumer (patient"s relative). A female patient (mother-in-law) of an unspecified age received unknown dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection; Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as dose number unknown, single for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient died from a blood clot to the brain on an unspecified date. The reporter personally think that it was from the Pfizer shot. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information about lot/batch number not available despite the follow-up attempts made. Follow-up attempts have been completed and no further information is expected.; Reported Cause(s) of Death: blood clot to the brain

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VAERS ID: 1655105 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac disorder, Condition aggravated, Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Heart disorder
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC202101091316

Write-up: 2 family"s members of hers died after receiving the Pfizer BioNtech covid 19 vaccine; One person who already had heart damage the vaccine exacerbated it and the spike proteins in their heart.; One person who already had heart damage the vaccine exacerbated it and the spike proteins in their heart.; This is a spontaneous report from a Non-contactable consumer. This consumer reported for 2 other patients that a patient of unspecified age and gender received bnt162b2 (BNT162B2), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose number unknown, single for covid-19 immunisation. Medical history included cardiac disorder from an unknown date and unknown if ongoing. The patient"s concomitant medications were not reported. The patient experienced 2 family"s members of hers died after receiving the pfizer biontech covid 19 vaccine on an unspecified date , one person who already had heart damage the vaccine exacerbated it and the spike proteins in their heart on an unspecified date with outcome of unknown. The patient died on an unspecified date. It was not reported if an autopsy was performed. The clinical course was reported as follows: Caller stated common knowledge that your vaccine and moderna create spike protein within the body, how do you as a company plan to fix that? It doesn"t stay in the arm, the spike protein is whats causing the excess problems. Caller reported 2 family"s members of hers died after receiving the Pfizer BioNtech covid 19 vaccine. One person who already had heart damage the vaccine exacerbated it and the spike proteins in their heart. She also reports various adverse events...damage to the heart , Guillen barre, women miscarrying their babies, other women starting periods after menopause. Caller stated that creator of the mRNA stated that its going to kill people within 3 years. Caller also states that white coat summit states its dangerous and unsafe. How you plan on helping theses people who have financial responsibility who can no longer work, care for their families, no longer here for their families or the Comirnaty vaccine damaged them unrecognizably. Caller states that the Comirnaty vaccine is not preventing or saving anybody from the covid 19 infection. No follow-up attempts are possible; information about lot/batch number cannot be obtained; Reported Cause(s) of Death: 2 family"s members of hers died after receiving the Pfizer BioNtech covid 19 vaccine

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VAERS ID: 1655107 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No

Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101091793

Write-up: a 19 year old female is dead from second dose of Pfizer Covid 19 vaccine; This is a spontaneous report from a Pfizer-sponsored program. This report reported similar events for two patients. This is the second of two reports. A female patient of 19-years-old received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE 2, SINGLE at the age of 19-years-old for covid-19 immunisation. The patient medical history and concomitant medications were not reported. A 19 years old female is dead from second dose of Pfizer Covid 19 vaccine. Both had no prior conditions with the heart and now are dead. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101091051 same report/drug/AE, different patients; Reported Cause(s) of Death: a 19 year old female is dead from second dose of Pfizer Covid 19 vaccine

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VAERS ID: 1655111 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 2021-04-19
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Dyspnoea, Renal disorder, Renal failure
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (narrow), Anaphylactic reaction (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), Acute central respiratory depression (broad), Pulmonary hypertension

(broad), Cardiomyopathy (broad), Chronic kidney disease (narrow), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
  Date died: 0000-00-00
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: Stroke (before 2020 so he was weak); Weakness
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101093322

Write-up: pass away/he died; couldn"t breathe; kidney dialysis fail; kidney stops; This is a spontaneous report from a contactable consumer (patient"s wife). A 73-year-old male patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 19Apr2021 as DOSE 2, SINGLE for COVID-19 immunization. Medical history included had a stroke year before 2020 so he was weak. Concomitant medications were not provided. The patient previously received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), on an unspecified date as single dose for COVID-19 immunization. The patient"s wife called on behalf of the patient seeking compensation due to the Pfizer-BioNTech Covid-19 Vaccine. It was stated patient 19Apr2021 they had the second vaccine together. And on Tuesday 1pm (in 2021) ambulance came in and got patient cause he couldn"t breathe, and they sent him to privacy hospital cause the kidney dialysis fail. They went in the morning 8"clock they go in there and they call 8:30 patient pass away in 2021. Patient had a stroke year before 2020 so he was weak they push him, and walk him, exercise and all kind of things. They didn"t warn that not supposed to have a second vaccine so patient and wife went no more a week later, 2 weeks, patient and wife had first and second shot and then patient went #. When patient died and his kidney stopped. Events reported as hospitalized in 2021. The outcome of the event pass away/he died was fatal, while of the other events was unknown. It was not reported if an autopsy was performed. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: pass away/he died

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VAERS ID: 1655112 (history)
Form: Version 2.0
Age:
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Chemotherapy (patient that was going through chemotherapy at age 80 for prostate cancer); Prostate cancer (patient that was going through chemotherapy at age 80 for prostate cancer.)
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101093359

Write-up: This person got the Covid 19 vaccine and died 3 days later; This is a spontaneous report from a contactable consumer. An 80-years-old male patient received dose of BNT162B2 (reported as COVID 19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunization. Medical history included prostate cancer and chemotherapy. The patient''s concomitant medications were not reported. The reporter stated she knew a radiology technician who mentioned the patient that was going through chemotherapy at age 80 for prostate cancer. This patient got the Covid 19 vaccine and died 3 days later. She thought it was a bad call for this patient to get the Covid 19 vaccine. She stated some people might think that because of being immune compromised those people might succumb to the Covid 19 virus. She asked in certain patient populations, like if someone is receiving chemotherapy, is it recommended that the Covid 19 vaccine be given. The report was serious with seriousness criteria-results in death. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for BNT162B2, was not provided and will be requested during follow up.; Reported Cause(s) of Death: This person got the Covid 19 vaccine and died 3 days later

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VAERS ID: 1655113 (history)
Form: Version 2.0
Age:

Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101093384

Write-up: 2 family''s members of hers died after receiving the Pfizer BioNtech covid 19 vaccine; This is a spontaneous report from a non-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 (Batch/Lot number was not reported) as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Reporter''s 2 family''s members of hers died after receiving the pfizer biontech covid 19 vaccine on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: 2 family''s members of hers died

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VAERS ID: 1655116 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101093930

Write-up: Passed away after being vaccinated; This is a spontaneous report from a Pfizer sponsored program from a Non-contactable consumer (reporter"s cousin). A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The reporter stated his cousin pass away after being vaccinated. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: Passed away after being vaccinated

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VAERS ID: 1655117 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

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

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Death, Drug ineffective, SARS-CoV-2 test
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: Covid; Test Result: Positive
CDC Split Type: USPFIZER INC202101094144

Write-up: Died due to the fact he received the Covid shot; Tested Positive for Covid after Vaccine; Tested Positive for Covid after Vaccine; This is a spontaneous report from a contactable consumer (patient"s friend). A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient"s medical history and concomitant medications were not reported. The consumer stated she knew that a friend (the patient) died due to the fact he received the Covid shot. The consumer stated all of the patient"s body and organs were spiked with proteins, and they were elevated from Covid. The patient was tested Positive for Covid after Vaccine on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the event tested Positive for Covid was unknown. The lot number for [bnt162b2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: died due to the fact he received the Covid shot

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VAERS ID: 1655118 (history)
Form: Version 2.0
Age: 33.0
Sex: Female
Location: Florida
Vaccinated: 2021-02-05
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-08-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	EN5318 / 1	LA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Hypoplastic left heart syndrome
SMQs:, Congenital, familial and genetic disorders (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-02-08
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: ASPIRIN [ACETYLSALICYLIC ACID]
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Cardiac operation NOS; Heart disorder
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101095126

Write-up: Hypoplastic left heart syndrome; This is a spontaneous report from a contactable consumer (patient"s father). A 33-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 05Feb2021 (reported as in the afternoon, after 1400) (Batch/Lot Number: EN5318) (at the age of 33-years-old) as DOSE 1, SINGLE for covid-19 immunisation. Medical history included heart issue with her heart from an unknown date, several heart surgeries from an unknown date. There were no additional vaccines administered on same date of the Pfizer Suspect. Prior Vaccinations (within 4 weeks) was none. There was no family medical history relevant to AE. Concomitant medication included ASPIRIN [acetylsalicylic acid] tablet taken for an unspecified indication, start and stop date were not reported. The consumer stated he did not keep close contact on all her medications since she was not living with her parents. The patient experienced hypoplastic left heart syndrome on an unspecified date and died for it on 08Feb2021. The consumer was reporting that his daughter passed away after she got the first Pfizer covid shot. She passed away in February. She got her shot on a Friday and passed away Monday evening. She did have a heart issue with her heart. At the time she had been through several heart surgeries and they did not realize the covid vaccine had issues for cardiac patients. The vaccine was recommended by her doctors. He reported she had hypoplastic left heart syndrome and that was the cause of death. She died on 08Feb2021. She died in her sleep, between 12:30am and 7:30am on 08Feb2021. She did not have an autopsy done at the time. He reported she had just got the shot and he was not sure if she died from natural causes or from the shot. In his heart he felt like she had been cut up enough and did not have an autopsy done. If he knew then the information that he knew now, then he would have had an autopsy done. She received vaccine basis that of her doctor"s recommendation and her mother"s panic about her covid and dyeing an ugly death, she had two cardiovascular surgeons that both

recommended to get the covid vaccine. There was no Emergency Room visit and no physician office visit. The patient died on 08Feb2021. An autopsy was not performed. The outcome of the event hypoplastic left heart syndrome was fatal.; Reported Cause(s) of Death: Hypoplastic left heart syndrome

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VAERS ID: 1655120 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-30

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

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

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101095685

Write-up: had a heart attack and died; This is a spontaneous report from a Pfizer-sponsored program from a contactable consumer (patient''s wife). A male patient of unknown age received first and second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), both on unknown date at single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient was fully vaccinated with the Pfizer Covid 19 vaccine, had a heart attack after both vaccine. The patient died due to heart attack. It was unknown if autopsy was done. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: had a heart attack and died.

VAERS ID: 1657901 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Ohio
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101097920

Write-up: death; This is a spontaneous report from a Pfizer-sponsored-program. A contactable consumer reported that a male patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on unspecified date at single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter stated "You are despicable. Your VAX caused my dad"s death. Apparently, lives are disposable as long as you make your money. I hope your family members don"t die slow, agonizing deaths from this vax like my dad did". It was unknown if autopsy was done. The outcome of the event was fatal. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: death

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VAERS ID: 1657902 (history)
Form: Version 2.0

Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101098146

Write-up: died; This is a spontaneous report from a contactable consumer. An 18-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot number and expiration date not provided) via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunisation. The patient"s medical history and concomitant medications were not reported. The patient died on an unspecified date due to unknown cause of death. Unknown treatment received. It was unknown if an autopsy was performed. The lot number for BNT162B2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: died

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VAERS ID: 1657906 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00

Submitted: 0000-00-00
Entered: 2021-08-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101100059

Write-up: lost a friend; This is a spontaneous report from a non-contactable consumer reporting same event under the same suspect product for 12 patients. This is one of 12 reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified routed of administration on unspecified date as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter had lost a friend. The cause of death was unknown. It was unknown if autopsy was done. The outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101100128 Same reporter/event/drug, different patient;US-PFIZER INC-202101100129 Same reporter/event/drug, different patient;US-PFIZER INC-202101100127 Same reporter/event/drug, different patient; Reported Cause(s) of Death: lost a friend

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VAERS ID: 1657907 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-08-31

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101100127

Write-up: I have lost 3 friends in Spain; This is a spontaneous report from a non-contactable consumer. This consumer reported similar events for twelve patients. This is one of twelve reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified routed of administration on unspecified date as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter stated "I have lost 7 friends in , 3 in , 1 in and a friend of my son 14 years." The cause of death was unknown. It was unknown if autopsy was done. The outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202101100059 Same reporter/event/product, different patient; Reported Cause(s) of Death: lost 3 friends

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VAERS ID: 1657909 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-08-31

Vaccination / Manufacturer	Lot /	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101100129

Write-up: lost a friend of my son 14 years; This is a spontaneous report from a non-contactable consumer reporting same event under the same suspect product for 12 patients. This is one of 12 reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified routed of administration on unspecified date as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter had lost a friend of the reporter''s son 14 years (seems the reporter''s son''s age who is not the patient). The cause of death was unknown. It was unknown if autopsy was done. The outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101100059 Same reporter/event/drug, different patient; Reported Cause(s) of Death: lost a friend of my son 14 years

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VAERS ID: 1660988 (history)
Form: Version 2.0
Age:
Sex: Male
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210860032

Write-up: DEATH; This spontaneous report received from a consumer via social media concerned a male of unspecified age, race and ethnicity. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry: unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date spike protein was found in every organ of the patient and patient died after 7 days of vaccination from an unknown cause. It was unknown whether autopsy was done or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210860032-COVID-19 VACCINE AD26.COV2.S-DEATH. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1661267 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Texas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

PFIZER/BIONTECH		

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101110314

Write-up: recently died from covid; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received bnt162b2 (COVID-19 VACCINE - MANUFACTURER UNKNOWN), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The reporter had a good friend who was fully vaccinated and didn''t know which vaccine he got, who recently died from covid. The reporter knew nothing was 100%. The patient died on an unspecified date in 2021. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: recently died from covid

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VAERS ID: 1661268 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Aneurysm
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101110596

Write-up: aneurysm; This is a spontaneous report from a non-contactable consumer. A patient of unspecified age and gender may received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. Caller claimed that some guy she knows got an aneurysm and died but she was not sure if that person took the vaccine. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: aneurysm

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VAERS ID: 1661269 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No

Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101110737

Write-up: passed away; This is a spontaneous report from a contactable consumer. A 51-year-old male patient received 1st dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as single dose for COVID-19 immunization. The patient"s medical history and concomitant medications were not reported. It was stated that patient passed away one week after receiving the 1st Pfizer covid vaccine injection. He was young and healthy. The patient died on an unspecified date due to unknown cause of death. It was unknown if an autopsy was performed. The lot number for BNT162B2, was not provided and will be requested during follow up.; Reported Cause(s) of Death: passed away

---

VAERS ID: 1661270 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No

Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101111871

Write-up: died at 36 years old.; This is a spontaneous report from a non-contactable consumer. A 36-years-old male patient received bnt162b2 (BNT162B2), on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient medical history and the concomitant medications were not reported. The patient died at 36 years old. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: died at 36 years old

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VAERS ID: 1661272 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebrovascular accident
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101113182

Write-up: stroke; This is a spontaneous report from a contactable consumer (patient"s brother). A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The reporter said that his brother passed away after receiving the vaccine and had a stroke on an unspecified date, and there was nothing wrong with him prior to that. The patient died on an unspecified date. It was not reported if an autopsy was performed. The reporter would like to know what exactly is in the shot and what the chemicals are that are in the shot. He has been recently notified that the FDA has approved the booster and wants to know what is in that one too. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up. ; Reported Cause(s) of Death: stroke

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VAERS ID: 1661274 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Virginia
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Alanine aminotransferase, Antibody test, Aspartate aminotransferase, Blood creatine phosphokinase MB, Blood fibrinogen, Blood lactic acid, Blood pressure measurement, C-reactive protein, Cardiac arrest, Catheterisation cardiac, Echocardiogram, Electrocardiogram, Fibrin D dimer, Haemoglobin, Heart rate, Interleukin level, International normalised ratio, Investigation, Myocarditis, Platelet count, Polymerase chain reaction, SARS-CoV-2 antibody test, Serum ferritin, Shock
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), Hypovolaemic shock conditions (narrow), Toxic-septic shock conditions (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypoglycaemic and neurogenic shock conditions (narrow), Acute central respiratory depression (broad), Cardiomyopathy (broad), Hypotonic-hyporesponsive episode (broad), Hypersensitivity (narrow), Respiratory failure (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad), COVID-19 (broad)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 0000-00-00
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:
Current Illness: Speech impairment NOS; Trisomy 21 (complicated by speech impairment)
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: alanine transferase; Result Unstructured Data: Test Result:7995 IU/l; Comments: u/l; Test Name: anti-nucleocapsid IgG; Test Result: Negative ; Test Name: aspartate transferase; Result Unstructured Data: Test Result:9003 IU/l; Comments: u/l; Test Name: creatine kinase myocardial band level; Result Unstructured Data: Test Result:252 ng/ml; Comments: normal value less than 5; Test Name: fibrinogen; Test Result: 100 mg/dl; Comments: dropped to 100 mg/dL; Test Name: Lactic acid; Result Unstructured Data: Test Result:6 mmol/L; Test Name: Lactic acid; Result Unstructured Data: Test Result:28 mmol/L; Comments: increased (NV - Less than 2); Test Name: blood pressure; Result Unstructured Data: Test Result:77/54 mmHg; Test Name: Cardiac catheterization; Result Unstructured Data: Test Result:showed no coronary obstructions; Test Name: CRP; Test Result: 13.1 mg/dl; Comments: highly elevated on admission; Test Name: Transthoracic echocardiogram; Test Result: 20 %; Comments: showed severe left ventricular systolic dysfunction (LVEF 20%) and a small circumferential pericardial effusion without tamponade; Test Name: electrocardiogram; Result Unstructured Data: Test Result:diffuse ST segment elevation; Comments: found to have diffuse ST segment elevation; Test Name: D-dimer; Result Unstructured Data: Test Result:4.21 ug/ml; Test Name: D-dimer; Result Unstructured Data: Test Result:greater than 20 ug/ml; Comments: increased critically high greater than 5.0; Test Name: hemoglobin; Result Unstructured Data: Test Result:11.5; Test Name: hemoglobin; Result Unstructured Data: Test Result:8.7; Comments: decreased; Test Name: heart rate; Result Unstructured Data: Test Result:133; Comments: per min; Test Name: Interleukin-6; Result Unstructured Data: Test Result:333 pg/mL; Comments: (NV less than 13); Test Name: INR; Result Unstructured Data: Test Result:2.0; Test Name: INR; Result Unstructured Data: Test Result:10.0; Comments: increased; Test Name: other respiratory virus; Test Result: Negative ; Comments: other common respiratory viruses more frequently associated with myocarditis was negative; Test Name: thrombocytes; Result Unstructured Data: Test Result:223 x10 9/l; Test Name: thrombocytes; Result Unstructured Data: Test Result:21 x10 9/l; Comments: developed progressively; Test Name: Polymerase chain reaction; Test Result: Negative ; Comments: for SARS-CoV2; Test Name: SARS-CoV2 NC igG; Test Result: Negative ; Test Name: SARS-CoV2 spike igG; Result Unstructured Data: Test Result:62.8; Comments: a.u./ml; Test Name: SARS-CoV2 spike protein IgG antibody; Test Result: Positive ; Comments: (62.8 arbitrary units/ml [NV less than 15.0]); Test Name: Ferritin; Result Unstructured Data: Test Result:23000 ng/ml; Comments: highly elevated on admission
CDC Split Type: USPFIZER INC202101116136

Write-up: fulminant myocarditis; recurrent cardiac arrest; refractory shock; This is a literature report from the International Journal of Cardiology, 2021, DOI:10.1016/j.ijcard.2021.08.018 entitled Fulminant myocarditis and systemic hyperinflammation temporally associated with BNT162b2 mRNA COVID-19 vaccination in two patients. This author reported similar events for two patients. This is first of two reports. A 27-year-old male with trisomy 21 complicated by speech impairment without history of cardiovascular disease presented in cardiogenic shock 2

days after his second vaccine dose. He had received the first dose without adverse effects. Approximately 36 h after the second dose, he developed nausea and vomiting. He presented to another hospital in shock (blood pressure 77/54 mmHg and heart rate 133/min) and found to have diffuse ST segment elevation in electrocardiogram (Fig. 1). Cardiac catheterization showed no coronary obstructions. Initially, creatine kinase myocardial band level (CK-MB) was 252 ng/mL (normal value [NV] < 5). Transthoracic echocardiogram showed severe left ventricular systolic dysfunction (LVEF 20%) and a small circumferential pericardial effusion without tamponade. A diagnosis of presumed fulminant pericarditis was made and methylprednisolone 1000 mg and human immunoglobulin (IVIG) 60 g were given. The course was complicated by hemodynamically unstable ventricular tachycardia refractory to electrical cardioversion followed by pulseless electrical activity. He was resuscitated with veno-arterial extracorporeal mechanical oxygenation (VA-ECMO). After return of circulation, he was supported by multiple vasopressors, mechanical ventilation, and renal replacement therapy (RRT). Despite these interventions, multiorgan failure and refractory shock persisted. Lactic acid increased from 6 to 28 mmol/L (NV less than 2), D-dimer increased from 4.21 to greater than 20 ug/mL (critically high $g5.0), INR increased from 2.0 to 10.0, and fibrinogen dropped to 100 mg/dL. C-reactive protein (CRP) and ferritin were highly elevated on admission at 13.1 mg/dL (NV less than 0.5) and 23,000 ng/mL, respectively, leading to a decision to administer anakinra (Kineret). Interleukin-6 level eventually came back highly elevated at 333 pg/mL (NV less than 13). thrombocytopenia developed progressively from 223x10 9 /L to 21x10 9 /L, while hemoglobin decreased from 11.5 to 8.7 g/dL. The patient demonstrated acute liver injury (ALI) with alanine transferase (ALT) and aspartate transferase (AST) at 7995 and 9003 u/L. Polymerase chain reaction (PCR) for SARS-CoV2 and other common respiratory viruses more frequently associated with myocarditis was negative. SARS-CoV2 spike protein IgG antibody was positive (62.8 arbitrary units/ml [NV less than 15), and anti-nucleocapsid IgG was negative, consistent with immunization status. Approximately 21 h after admission, patient died due to recurrent cardiac arrest and refractory shock. Family declined request for autopsy. In summary, both cases presented features of fulminant myocarditis with a temporal association with the BNT162b2 mRNA Covid-19 vaccination, in absence of other apparent causes and with unique features of systemic hyperinflammation associated with refractory shock. These cases identify the need for awareness of a potential, albeit extremely rare, link of BNT162b2 mRNA Covid-19 vaccination associated with fulminant myocarditis as part of a severe systemic hyperinflammatory syndrome, requiring mechanical cardiac support and, most importantly, immunosuppressive therapy. The degree of the inflammatory biomarkers and multiorgan dysfunction seen in the two cases described, as in some patients with Covid-19, is indeed out of proportion to with hemodynamic failure and shock and it is not rapidly resolved by cardiac mechanical support, reflecting a significant degree of vasoplegia. The optimal immunosuppressive treatment for systemic hyperinflammation associated with fulminant myocarditis is unknown. They chose a combination of high dose methylprednisolone, IVIG and anakinra (Table 1) as a strategy used across a variety of immunologic and rheumatologic diseases characterized with inappropriate macrophage activation. Table 1 Key biomarkers at admission and immunomodulating therapy: Patient #1: Presentation included Symptoms: Nausea and vomiting. Vaccine type was BNT162b2 mRNA Covid-19. Timing was 2 days after second dose. Hemodynamics included Hypotension and tachycardia. ECG: ST segment elevation. Echocardiogram: LVEF 20%. Biomarkers included: CRP: 13.1 mg/dl; Interleukin-6 : 333 pg/ml; Ferritin : 23,000 ng/ml; INR : 2.0; D-dimer : 4.21 ug/ml; Lactic acid : 6 mmol/l; CK-MB : 252 ng/ml; SARS-CoV2 spike IgG : 62.8 a.u./ml; SARS-CoV2 NC IgG : negative. Immunosuppressive Therapy included: Methylprednisolone : 1000 mg; Immunoglobulins : 60 mg; Anakinra : 100 mg (2 doses). Outcome was Deceased 21 h after admission. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender"s Comments: Based on temporal association and known drug profile, a contributory role of the suspect drug cannot be excluded in the development of events myocarditis, cardiac arrest, and shock. This case will be reassessed upon receipt of additional information.,Linked Report(s) :PFIZER INC-

202101116531 Same article/ drug/event and different patient; Reported Cause(s) of Death: recurrent cardiac arrest; refractory shock; fulminant myocarditis

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VAERS ID: 1661277 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-01

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101127070

Write-up: she died; This is a spontaneous report from a contactable consumer (patient was reporter"s girlfriend). A 49-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient experienced she died (death, medically significant) on an unspecified date immediately, 49-year-old from Pfizer. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for BNT162B2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: she died

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VAERS ID: 1665394 (history)

Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-02

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, SARS-CoV-2 test, Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Parkinson''s disease
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Result Unstructured Data: Negative
CDC Split Type: USJNJFOC20210855586

Write-up: DEATH; SUSPECTED CLINICAL VACCINE FAILURE; SUSPECTED COVID-19 INFECTION; This spontaneous report received from a consumer via social media concerned an 87 year old male. The patient''s height, and weight were not reported. The patient''s concurrent conditions included Parkinson''s disease. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown)1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient was fully vaccinated still got covid-19 (suspected covid-19 infection and suspected clinical vaccine failure). Patient was finally tested negative for covid-19. It was too much for patient''s Parkinson body he died in his sleep. The cause of death was unknown. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of suspected covid-19 infection and suspected clinical vaccine failure was not reported. This report was serious (Death).; Sender''s Comments: V0: 20210855586-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210855586-COVID-19 VACCINE AD26.COV2.S-Suspected clinical vaccination failure. This event(s) is considered not related. The event(s) has a

compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1665438 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: California
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-02

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210860177

Write-up: DEATH; This spontaneous report received from a consumer via social media (medical journal) concerned multiple patients of unknown race and ethnicity. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: UNKNOWN) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. The reporter read in journal that 22 people died after receiving the vaccine. The cause of death was unknown. It was unknown if autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210860032.; Sender"s Comments: V0: 20210860177-Covid-19 vaccine ad26.cov2.s -Death. This event(s) is considered unassessable. The event(s) has a

compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1665632 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Arkansas
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-02

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Dermatosis
SMQs:, Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: died; Dermatosis; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (died) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced DEATH (died) (seriousness criteria death and medically significant) and DERMATOSIS (Dermatosis). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, DERMATOSIS (Dermatosis) outcome was unknown. Concomitant medication and treatment drug were not reported. Company Comment: Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Sender''s Comments: Very limited information regarding this event/s has been provided at this time. Further information has been

requested.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1669558 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-03

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, 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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210855833

Write-up: DEATH; SEIZURES; This spontaneous report received from a consumer via a company representative via social media (twitter) concerned a male of an unspecified age. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient developed seizure and then died. As per the reporter, "I know an hour after taking the vaccine this 100 percent healthy kid had seizures. Next thing the patient was dead." It was unknown if an autopsy was performed or not. The cause of death was not reported. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died on an unspecified date, and the outcome of

seizures was not reported. This report was serious (Death).; Sender''s Comments: V0: 20210855833-covid-19 vaccine ad26.cov2.s -Death, Seizures. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1669567 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-03

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

Administered by: Other **Purchased by:** ?
Symptoms: Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-24
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? Yes
Hospitalized? Yes, 7 days
Extended hospital stay? No
Previous Vaccinations:
Other Medications:
Current Illness: Atrial fibrillation; Diabetes (Controlled); Hypertension
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210858457

Write-up: SUSPECTED COVID-19 INFECTION; SUSPECTED CLINICAL VACCINATION FAILURE; This spontaneous report received from a consumer concerned a 94 year old male and unspecified race & ethinicity. Initial information was processed along with the additional information received on 27-AUG-2021 The patient''s height, and weight were not reported. The patient''s concurrent conditions included: diabetes, hypertension, and atrial fibrillation. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) 1 in total dose was administered on APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. The consumer reported that father was in the

emergency room on 17-AUG-2021 and was hospitalized. On 18-AUG-2021, the patient was on ventilator and died on 24-AUG-2021 with the delta variant. On an unspecified date, the patient had suspected covid-19 infection, and suspected clinical vaccination failure. On 24-AUG-2021, the patient died from covid-19 infection. The patient was hospitalized for 7 days. It was unspecified if an autopsy was performed The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of suspected covid-19 infection on 24-AUG-2021. This report was serious (Death, and Hospitalization Caused / Prolonged).; Sender"s Comments: V0: 20210858457-covid-19 vaccine ad26.cov2.s- suspected clinical vaccination failure. This event is considered not related. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event than the drug. Specifically: SPECIAL SITUATIONS 20210858457-covid-19 vaccine ad26.cov2.s- suspected covid-19 infection. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: COVID-19 INFECTION

VAERS ID: 1669577 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-03

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210860208

Write-up: DEATH; This spontaneous report received from a consumer via company representative concerned 726 patients of unspecified sex, race and ethnic origin. The patients height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin was not reported, batch number: unknown and expiry date: unknown) dose was not reported, 1 total administered, start therapy date was not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patients died from unknown cause of death. It was unknown that whether autopsy performed for reported patients. This report was serious (Death).; Sender"s Comments: V0: 20210860208-COVID-19 VACCINE AD26.COV2.S-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1669586 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-03

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

Administered by: Other **Purchased by:** ?
Symptoms: Cerebrovascular accident
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)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-19
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Comments: Unknown
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210900460

Write-up: TWO STROKES; This spontaneous report received from a consumer via a company representative through social media concerned a female of unspecified age, race and ethnicity. The patient''s weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. It was reported that, on an unspecified date, the vaccine caused blood clotting throughout the patient''s body resulting in two strokes and ultimately her death on 19-AUG-2021. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. On 19-AUG-2021, the patient died from two strokes. This report was serious (Death).; Sender''s Comments: V0: 20210900460 - JANSSEN COVID-19 VACCINE Ad26.COV2.S- Two strokes. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: TWO STROKES

VAERS ID: 1672341 (history)
Form: Version 2.0
Age: 80.0
Sex: Male
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Blister, Death
SMQs:, Severe cutaneous adverse reactions (broad), Hypersensitivity (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: passed away; bubbles like blisters in the face/neck /back /arms / legs; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (passed away) and BLISTER (bubbles like blisters in the face/neck /back /arms /legs) in an 80-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced DEATH (passed away) (seriousness criteria death and medically significant) and BLISTER (bubbles like blisters in the face/neck /back /arms / legs) (seriousness criterion death). The reported cause of death was bubbles like blisters in the face/neck /back /arms / legs. It is unknown if an autopsy was performed. Concomitant products were not provided. Treatment medication were not reported. Patient was healthy in his 80s. The patient had bubbles like blisters in the face and in the back of the neck, back and arms, and legs. This is a case of death in a 80-year-old male patient after receiving vaccine. The patient also experienced blisters in the face and in the back of the neck, back and arms, and legs. Very limited information has been provided at this time.; Sender''s Comments: This is a case of death in a 80-year-old male patient after receiving vaccine. The patient also experienced blisters in the face and in the back of the neck, back and arms, and legs.Very limited information has been provided at this time.; Reported Cause(s) of Death: passed away; bubbles like blisters in the face/neck /back /arms / legs

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VAERS ID: 1673094 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (MODERNA)) / MODERNA	UNKNOWN / 1	- / OT

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiomegaly, Death, Palpitations, Product administered to patient of inappropriate age
SMQs:, Cardiac failure (broad), Arrhythmia related investigations, signs and symptoms (broad), Cardiomyopathy (broad), Medication errors (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:

Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: death; teenager got palpitation; her heart enlarged; teenager got a covid-19 vaccine.; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (death), PALPITATIONS (teenager got palpitation), CARDIOMEGALY (her heart enlarged) and PRODUCT ADMINISTERED TO PATIENT OF INAPPROPRIATE AGE (teenager got a covid-19 vaccine.) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced DEATH (death) (seriousness criteria death and medically significant), PALPITATIONS (teenager got palpitation) (seriousness criterion death), CARDIOMEGALY (her heart enlarged) (seriousness criterion death) and PRODUCT ADMINISTERED TO PATIENT OF INAPPROPRIATE AGE (teenager got a covid-19 vaccine.) (seriousness criterion death). The reported cause of death was teenager got palpitation, her heart enlarged and teenager got a covid-19 vaccine.. It is unknown if an autopsy was performed. Concomitant product use was not provided by the reporter. No treatment information was provided. Very limited information regarding these events has been provided at this time. Further information has been requested.; Sender''s Comments: Very limited information regarding these events has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: teenager got palpitation; her heart enlarged; teenager got a covid-19 vaccine.

VAERS ID: 1673406 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Infection
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No

ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101109686

Write-up: breakthrough infection and died; 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 (Batch/Lot number was not reported) as dose number unknown, single for COVID-19 immunisation. The patient''s medical history and concomitant medications were not reported. The consumer stated that on the website, it says that it''s not FDA approved, as well as voicemails. She went on to say that it only takes a minute to update those things, and she would think with something this large, they would have been ready. She stated that she was aware that the VAERS website contains side effects and things that have happened. She stated that other vaccines have been halted after 1 to 2 deaths. She is aware of a couple of people dying as a direct result of the Pfizer BioNTech vaccine, as well as as friend that had a breakthrough infection and died on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for BNT162b2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: breakthrough infections and has died

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VAERS ID: 1673422 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No

Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101113082

Write-up: Died after taking the vaccine; This is a spontaneous report from a non-contactable consumer. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date as single dose for COVID-19 immunization. Medical history and concomitant medications were Not Provided. The reporter said his cousin died after taking the Covid vaccine and reported as serious due to death. The caller mentioned about that our website or recorded line doesn"t say FDA approval and she told him that will be updated by the end of the day and they are aware of that. Reporter then started to say is the president lying and the agent ended the call at that time. Investigation Assessment was Not Provided. It was not reported if Autopsy Done. The outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: died after taking the vaccine

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VAERS ID: 1673454 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101123341

Write-up: passed away; This is a spontaneous report from a Pfizer-sponsored program via Regulatory Authority Support received from a non-contactable consumer. The consumer reported same events for two patients (two friends). This is one of the two reports. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE 2, SINGLE, dose 1 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE 1, SINGLE for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient took both doses and passed away on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101123407 Same reporter/suspect product/event, different patients; Reported Cause(s) of Death: took both doses and passed away

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VAERS ID: 1673460 (history)
Form: Version 2.0
Age: 94.0
Sex: Female
Location: Texas
Vaccinated: 2021-03-16
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	LA / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Renal failure
SMQs:, Rhabdomyolysis/myopathy (broad), Acute renal failure (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), Chronic kidney disease (narrow), Tumour lysis syndrome (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad)

Life Threatening? No
Birth Defect? No
Died? Yes

Date died: 2021-05-18
Permanent Disability? No
Recovered? No
Office Visit? Yes
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: Temazepam; Centrum Silver [Ascorbic Acid; Calcium; Minerals Nos; Retinol; Tocopheryl Acetate; Vitamin B NOS; Vitamins; Ocuvite [Ascorbic Acid; Beta carotene; Copper; Tocofersolan (TPGS); Zinc]
Current Illness:
Preexisting Conditions: Comments: List of non-encoded Patient Relevant History: Patient Other Relevant History 1: None, Comment: other medical history: none, she was just old.
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101127960

Write-up: Renal Failure; This is a spontaneous report from a contactable consumer. A 94-year-old female patient (not pregnant) received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 2 via an unspecified route of administration, administered in Arm Left on 16Mar2021 15:00 (Batch/Lot number was not reported) (at the age of 94-year-old, not pregnant) as single dose for COVID-19 immunisation. Medical history: none, she was just old. Concomitant medications included temazepam; ascorbic acid, calcium, minerals nos, retinol, tocopheryl acetate, vitamin b nos, vitamins nos, zinc (CENTRUM SILVER); ascorbic acid, betacarotene, copper, tocofersolan, zinc (OCUVITE). The patient previously received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE) dose 1 on 23Feb2021 at 12:00 PM in left arm at the age of 94-year-old for COVID-19 immunization and experienced Renal Failure. No covid prior vaccination. No other vaccine in four weeks. No covid tested post vaccination. No known allergies. The patient experienced periodic uncontrollable shaking renal failure. AE resulted in: Doctor or other healthcare professional office/clinic visit. No treatment. The patient died on 18May2021. An autopsy was not performed. The lot number for [BNT162B2], was not provided and will be requested during follow up.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101102004 same reporter, patient, product, event, different doses and potential; Reported Cause(s) of Death: Renal Failure

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VAERS ID: 1673464 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

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

Administered by: Unknown **Purchased by:** ?

Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101129798

Write-up: who got the vaccine and died three days later; This is a spontaneous report from a non-contactable other hcp. The reporter reported for herself and another patient. This was for another patient. A patient of unspecified age and gender received bnt162b2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient got the vaccine and died three days later on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender"s Comments: The causal relationship between bnt162b2 and the event death cannot be excluded as 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.; Reported Cause(s) of Death: who got the vaccine and died three days later

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VAERS ID: 1673473 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

Vaccination / Manufacturer	Lot / Dose	Site / Route

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101134213

Write-up: died; This is a spontaneous report from a contactable consumer. This consumer reported for an adult male patient. An adult male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection. Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as DOSE NUMBER UNKNOWN, SINGLE for COVID-19 immunization. Reason for no lot number of Pfizer COVID-19 Vaccine: complainant hung up abruptly/transfer incomplete. The patient medical and concomitant medication history were reported. Other products, patient history and investigation assessment not provided. The patient experienced died on an unspecified date. It was not reported if an autopsy was performed. Caller was asking why Pfizer not reporting the side effects to VAERS? Caller states that the data on VAERS has stopped counting the fatalities from the Pfizer COVID-19 vaccine and the death count exceeds 12,000. Caller states that the Pfizer COVID-19 vaccine has metals and was asking how was it being passed through the FDA and Pfizer? Normally with other shots, they take years to get approved. FDA only allows, if people die, they did not put it on the market. Caller states that the data on VAERS has stopped counting the fatalities from the Pfizer COVID-19 vaccine and the death count exceeds 12,000. Caller asked if we had COVID, we had the antibodies, why do we have to get the shot. The lot number was not provided. Information on lot number has been requested during follow up.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202101134188 same reporter/drug/event, different patient.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1673475 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-04

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101134490

Write-up: I have lost 3 friends in Spain; This is a spontaneous report from a non-contactable consumer reporting same event under the same suspect product for 12 patients. This is one of 12 reports. A patient of unspecified age and gender received BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number not reported) via an unspecified routed of administration on unspecified date at a single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter had lost a friend in Spain. The cause of death was unknown. It was unknown whether an autopsy was done. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender"s Comments: Linked Report(s) : US-PFIZER INC-202101100059 same reporter/drug/event, different patient; Reported Cause(s) of Death: I have lost 3 friends in Spain

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VAERS ID: 1677072 (history)
Form: Version 2.0
Age: 20.0
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00

Entered: 2021-09-07

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210902754

Write-up: DIED; This spontaneous report received from consumer social media via a company representative concerned a 20 year old female of unknown race and ethnicity. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine (suspension for injection, route of admin not reported, batch number: Unknown, expiry: unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient died from unknown cause of death and it was unknown if an autopsy was performed. The action taken with covid-19 vaccine was not applicable. This report was serious (Death). This case, from the same reporter is linked to 20210902775.; Sender"s Comments: V0:20210902754-covid-19 vaccine Died. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1677074 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-07

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

Administered by: Other **Purchased by:** ?
Symptoms: COVID-19
SMQs:, Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210903589

Write-up: COVID-19; This spontaneous report received from a company representative concerned multiple patients of unknown race and ethnic origin. No past medical history or concurrent conditions were reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, start therapy date were not reported, 1 total for prophylactic vaccination. The batch number was not reported. Per procedure no follow up will be requested for this case. No concomitant medications were reported. It was reported that "529 people had died of COVID-19 since 01-APR-2021. According to the ADPH, this included twenty people who were vaccinated, who did their part in the fight against COVID-19". It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210903589- JANSSEN COVID-19 VACCINE Ad26.COV2.S- COVID-19. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: COVID-19

VAERS ID: 1677280 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Illness
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: unknown cause of death; got real sick; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of DEATH (unknown cause of death) and ILLNESS (got real sick) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced DEATH (unknown cause of death) (seriousness criteria death and medically significant) and ILLNESS (got real sick) (seriousness criterion death). The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant medication information not provided. Treatment information not provided. Reporter girl friend is a truck driver and knows a few people who got real sick from getting it and passed away. He said they got the Moderna vaccine. He does not know if it is from the shot. He did not provide any further information. Very limited information regarding the patient, product, events, clinical details and concomitant medications has been provided at this time. Cause of death not provided. Lack of information limits the causality assessment.; Sender''s Comments: Very limited information regarding the patient, product, events, clinical details and concomitant medications has been provided at this time. Cause of death not provided. Lack of information limits the causality assessment.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1677583 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Myocarditis
SMQs:, Cardiomyopathy (broad), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101134355

Write-up: she knew/heard someone past away (die) due to inflammation of the heart; This is a spontaneous report from a contactable consumer via Pfizer Sponsored Program. This consumer reported for two patients. This is the second of two reports. A female patient of an unspecified age received BNT162B2, via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as DOSE NUMBER UNKNOWN, SINGLE for COVID-19 immunization. The patient''s medical history and concomitant medications were not reported. The reporter would like to know if inflammation of the heart was a commonly reported side effect post-vaccination of the COMIRNATY vaccine and if there had been a lot of reports about it. The reporter stated that she knew/heard someone past away (die) due to inflammation of the heart (onset date not reported) as an AE post-vaccination. The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Sender''s Comments: Linked Report(s) :PFIZER INC-202101132810 Same reporter/drug, different patient/AE; Reported Cause(s) of Death: she knew/heard someone past away (die) due to inflammation of the heart

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VAERS ID: 1677599 (history)
Form: Version 2.0
Age:
Sex: Male

Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-07

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Cerebral thrombosis
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? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions: Medical History/Concurrent Conditions: Hashimoto''s disease
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101139821

Write-up: Got a blood clot that went to his brain and died; This is a spontaneous report from a contactable consumer (patient''s friend) based on information received by Pfizer, on behalf of Moderna (Case: MOD21-136450). A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Lot number and expiration date was not reported) as dose 2, single for COVID-19 immunization. Medical history included Hashimoto''s disease. The patient''s concomitant medications were none. Historical vaccine included BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Lot number and expiration date was not reported) as dose 1, single for COVID-19 immunization. On an unspecified date, the patient got a blood clot that went to his brain and had died on an unspecified date after 16-18 days of being fully vaccinated. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Reported Cause(s) of Death: Got a blood clot that went to his brain and died

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VAERS ID: 1678243 (history)
Form: Version 2.0
Age:

Sex: Female
Location: Arizona
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-07

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

Administered by: Other **Purchased by:** ?
Symptoms: Thrombosis
SMQs:, Embolic and thrombotic events, vessel type unspecified and mixed arterial and venous (narrow), Thrombophlebitis (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210902860

Write-up: BLOOD CLOTS; This spontaneous report received from a consumer via a company representative from social media (Twitter) concerned multiple (two female) patients. The patient"s weight, height, and medical histories were not reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, start therapy date was not reported for prophylactic vaccination. Batch numbers were not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, multiple (two) patients developed blood clots. As per the reporter, "We already know JnJ had to halt Vax distribution temporarily because of 2 reported deaths from blood clots in women". Patients died on unspecified date(s). It was not reported if an autopsy was performed. Action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0:20210902860- covid-19 vaccine ad26.cov2.s-Blood Clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOTS

VAERS ID: 1678295 (history)
Form: Version 2.0

Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-07

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20213

Write-up: Mother passed away; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (Mother passed away) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. Death occurred on an unknown date The cause of death was not reported. It is unknown if an autopsy was performed. Concomitant product use was not provided by the reporter. Treatment information was not provided. Reporter did not allow further contact; Sender''s Comments: Very limited information regarding this event has been provided at this time. Further information has been requested to confirm if this is a valid case.; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1681562 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00

Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Dementia, Helplessness, Mobility decreased
SMQs:, Dementia (narrow), Parkinson-like events (broad), Noninfectious encephalopathy/delirium (broad), Depression (excl suicide and self injury) (narrow), Tendinopathies and ligament disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? Yes
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness: Memory disturbance (patient had minor memory problems)
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USMODERNATX, INC.MOD20212

Write-up: it lead to the death of my mother; her mind went totally of the rails,She went from some minor memory problems to full blowing dementia/she became with dementia; she became totally helpless; unfunctioning after the second shot; This spontaneous case was reported by a consumer and describes the occurrence of DEATH (it lead to the death of my mother), DEMENTIA (her mind went totally of the rails,She went from some minor memory problems to full blowing dementia/she became with dementia), HELPLESSNESS (she became totally helpless) and MOBILITY DECREASED (unfunctioning after the second shot) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. Concurrent medical conditions included Memory disturbance (patient had minor memory problems). On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced DEATH (it lead to the death of my mother) (seriousness criteria death and medically significant), DEMENTIA (her mind went totally of the rails,She went from some minor memory problems to full blowing dementia/she became with dementia) (seriousness criteria disability and medically significant), HELPLESSNESS (she became totally helpless) (seriousness criterion disability) and MOBILITY DECREASED (unfunctioning after the second shot) (seriousness criterion disability). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, DEMENTIA (her mind went totally of the rails,She went from some minor memory problems to full blowing dementia/she became with dementia), HELPLESSNESS (she became totally helpless) and MOBILITY DECREASED (unfunctioning after the second shot) outcome was unknown. No concomitant medications or treatment details were reported. The patient was in

her 90s and was reportedly a vibrant person in full health and fully functioning. She just got her driving license and was driving, cooking, cleaning and walking. But within a week from the second shot she developed dementia and lead to her death. This a report of dead after receiving the product in a 90-years old female patient with Relevant Medical History of Minor memory problems.It is unknown if an autopsy was performed. Based on the current available information and temporal association between the use of the product, and the start date of the events, a causal relationship cannot be excluded. Very limited information regarding the event has been provided. Further information has been requested . This case was linked to MOD-2021-294328 (Patient Link).; Sender''s Comments: This a report of dead after receiving the product in a 90-years old female patient with Relevant Medical History of Minor memory problems.It is unknown if an autopsy was performed. Based on the current available information and temporal association between the use of the product, and the start date of the events, a causal relationship cannot be excluded. Very limited information regarding the event has been provided. Further information has been requested .; Reported Cause(s) of Death: Unknown cause of death

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VAERS ID: 1682014 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (PFIZER-BIONTECH)) / PFIZER/BIONTECH	- / 2	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:

CDC Split Type: USPFIZER INC202101123407

Write-up: passed away; This is a spontaneous report received from a non-contactable consumer from a Pfizer-sponsored program. The consumer reported same events for two patients (two friends). This is one of the two reports. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on an unspecified date as dose 2, single; dose 1 via an unspecified route of administration on an unspecified date as dose 1, single for covid-19 immunisation. The patient''s medical history and concomitant medications were not reported. The patient took both doses and passed away on an unspecified date. The patient died on an unspecified date. It was not reported if an autopsy was performed. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101123341 Same reporter/drug/event, different patient; Reported Cause(s) of Death: passed away

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VAERS ID: 1682056 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Cardiac arrest, Myocarditis
SMQs:, Torsade de pointes/QT prolongation (broad), Anaphylactic reaction (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), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Immune-mediated/autoimmune disorders (broad)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:

Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101138389

Write-up: Died of cardiac arrest; Inflammation on his heart; This is a spontaneous report from a contactable consumer or other non-HCP (patient friend) via Medical Information Team. A male patient of an unspecified age received BNT162b2 (PFIZER BIONTECH COVID-19 mRNA VACCINE, Solution for injection, Lot number- UNKNOWN) via an unspecified route of administration as dose number unknown single for COVID-19 immunization. Patient medical history and concomitant medications were not reported. Patient died recently and he was a bigger man. He experienced inflammation on his heart and died of cardiac arrest. It was unknown if an autopsy was done. The outcome of the event inflammation on his heart was unknown. The lot number for [BNT162B2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: Died of cardiac arrest

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VAERS ID: 1682092 (history)
Form: Version 2.0
Age:
Sex: Male
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death, Drug ineffective
SMQs:, Lack of efficacy/effect (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101139471

Write-up: My dad has been on Pfizer vaccine and he passed away from the complications because of it; He passed away, because of your vaccine; your vaccine, that doesn"t work; This is a spontaneous report from a contactable consumer or other non hcp. A male patient of an unspecified age received bnt162b2 (BNT162B2, Formulation: Solution for injection, Lot number was not reported, Expiry date: Unknown) via an unspecified route of administration on an unspecified date as dose number unknown, single for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. It was reported that on an unspecified date, my dad has been on Pfizer vaccine and he passed away from the complications because of it, he passed away, because of your vaccine. Reporter stated, he passed away, because of your vaccine. He never went to other house. Nobody came to his house. Last time he left the house when he got the vaccine and he died from it. Reporter want all to hold up to your vaccine because your vaccine, that doesn"t work. The patient died on an unspecified date. It was not reported if an autopsy was performed. The outcome of the event was fatal. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Reported Cause(s) of Death: My dad has been on Pfizer vaccine and he passed away from the complications because of it; He passed away, because of your vaccine

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VAERS ID: 1682095 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Utah
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:

Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101141821

Write-up: my friend"s grandmother died; This is a spontaneous report from a contactable consumer reported for friend"s grandmother via a female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date (Lot number was not reported) as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The reporter stated that her friend"s grandmother died with the Pfizer COVID 19 vaccine on an unspecified date. It was not reported if an autopsy was performed. The lot number for BNT162B2 was not provided and will be requested during follow up.; Reported Cause(s) of Death: my friend"s grandmother died

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VAERS ID: 1683561 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

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

Administered by: Other **Purchased by:** ?
Symptoms: Death, Hospitalisation, Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: USJNJFOC20210910577

Write-up: HOSPITALISATION; SUSPECTED CLINICAL VACCINATION FAILURE; SUSPECTED COVID-19 INFECTION; DEATH; This spontaneous report received from social

media via a company representative concerned multiple patients. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported,1 total for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the 15,739 patients got COVID-19 among people who were fully vaccinated (suspected covid-19 infection, suspected clinical vaccination failure), Of those, 571 had been hospitalised (for unspecified days), and 131 had died. The patients died due to unknown reason on an unspecified date. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. Outcome of suspected covid-19 infection, hospitalisation and suspected clinical vaccination failure was not reported. This report was serious (Death, Hospitalization Caused / Prolonged, and Other Medically Important Condition). This case, from the same reporter is linked to 20210902916, 20210909398 and 20210909281. This case was associated with Product Quality Complaint.; Sender''s Comments: V0; 20210910577-covid-19 vaccine ad26.cov2. s-Death, Hospitalization. These events are considered unassessable. The events have a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events. V0: - 20210910577- covid-19 vaccine ad26.cov2 - Suspected clinical vaccination failure. This event is considered not related. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event than the drug. Specifically: SPECIAL SITUATIONS.; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

VAERS ID: 1683731 (history)
Form: Version 2.0
Age:
Sex: Female
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-08

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

Administered by: Unknown **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No

Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USPFIZER INC202101134188

Write-up: female in her 40s that died after getting the Pfizer covid vaccine; This is a spontaneous report from a contactable consumer. This consumer reported for a female patient in her 40''s which included that. A 5-decade-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Batch/Lot number was not reported), via an unspecified route of administration on an unspecified date as dose number unknown, single for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Other Products, Patient History and Investigation Assessment was Not Provided. The patient experienced female in her 40s that died after getting the pfizer covid vaccine (death) (death, medically significant) on an unspecified date. Reporter reports that three people got the Pfizer covid 19 vaccine. One is a 63-year-old female, and she had a stroke her verbatim was bleed in the head. Then a female in her 40s that died after getting the Pfizer covid vaccine. Then a male that was 25-30 years old and he died. She does not have further details to provide on female in her 40s that died after the covid vaccine. Later Caller stated during call that she was speaking with another agent, but it was only this Caller on the line when this agent answered. Caller states that she knows a 63-year-old female who received the Pfizer Covid-19 vaccine then had a stroke, "bleed in the head" afterwards. Caller also states that she knew a female in a 40''s (years) and a male between the age of 25-30 years who both died after receiving the Pfizer Covid-19 vaccine. Caller is asking why is Pfizer not reporting the side effects to VAERS? Caller states that the data on VAERS has stopped counting the fatalities from the Pfizer Covid-19 vaccine and the death count exceeds 12,000. Caller stated that the agent before this agent was unable to answer any of her questions (this agent unaware of which Pfizer department due to no warm transfer being performed and the Caller was the only person on the line when this agent answered the Call). Caller provided the following possible feedback: Caller asks why did withheld, approve the Pfizer Covid- 19 vaccine? Caller asks why does she have anything to do with this approval of the Pfizer Covid-19 vaccine? Caller states that the Pfizer Covid-19 vaccine has metals and is asking how is it being passed through the FDA and Pfizer? Normally with other shots, they take years to get approved. FDA only allows, if people die, they don''t put it on the market. Caller states that the data on VAERS has stopped counting the fatalities from the Pfizer Covid-19 vaccine and the death count exceeds 12,000. Caller is asking if "We had COVID, we had the antibodies, why do we have to get the shot?" Confirmed with caller that she is talking about mandates from withheld and employers. Caller asks why did withheld, approve the Pfizer Covid-19 vaccine Caller asks why does she have anything to do with this approval of the Pfizer Covid-19 vaccine Caller states that the Pfizer Covid-19 vaccine has metals and is asking why is it being approved? The patient died on an unspecified date. It was not reported if an autopsy was performed. The lot number for the vaccine, [BNT162B2], was not provided and will be requested during follow up.; Sender''s Comments: Linked Report(s) : US-PFIZER INC-202101134213 same reporter/drug/event, different patient.; Reported Cause(s) of Death: unknown cause of death

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VAERS ID: 1685448 (history)
Form: Version 2.0

Age: 83.0
Sex: Female
Location: Florida
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-09

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

Administered by: Senior Living **Purchased by:** ?
Symptoms: Acute respiratory failure, COVID-19, Death, Dyspnoea, Fatigue, Gait inability, Laboratory test, Mobility decreased, SARS-CoV-2 test positive, Septic shock
SMQs:, Anaphylactic reaction (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 (narrow), Anaphylactic/anaphylactoid shock conditions (broad), Hypoglycaemic and neurogenic shock conditions (broad), Dystonia (broad), Parkinson-like events (broad), Acute central respiratory depression (narrow), Pulmonary hypertension (broad), Guillain-Barre syndrome (broad), Cardiomyopathy (broad), Hypersensitivity (broad), Respiratory failure (narrow), Tendinopathies and ligament disorders (broad), Infective pneumonia (broad), Sepsis (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-27
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications: None
Current Illness: None
Preexisting Conditions: Alzheimer''s
Allergies: None
Diagnostic Lab Data: Various Test Ran
CDC Split Type:

Write-up: Tiredness unable to get out of bed and walk Acute Respiratory Failure COVID 19 Positive Septic Shock Shortness of Breath Expired on 08/27/2021 *She lived with a caregiver. Her son called to report but had limited information. He says that the caregiver has more information.

VAERS ID: 1689728 (history)
Form: Version 2.0
Age: 63.0

Sex: Male
Location: Unknown
Vaccinated: 2021-03-30
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-10

Vaccination / Manufacturer	Lot / Dose	Site / Route
COVID19: COVID19 (COVID19 (JANSSEN)) / JANSSEN	1802068 / UNK	- / -

Administered by: Unknown **Purchased by:** ?
Symptoms: COVID-19, Condition aggravated, Cough, Death, End stage renal disease, Intensive care, Pyrexia, SARS-CoV-2 test positive, Septic cerebral embolism, Shock, Staphylococcal bacteraemia
SMQs:, Rhabdomyolysis/myopathy (broad), Anaphylactic reaction (narrow), Neuroleptic malignant syndrome (broad), Anticholinergic syndrome (broad), Shock-associated circulatory or cardiac conditions (excl torsade de pointes) (narrow), Hypovolaemic shock conditions (narrow), Toxic-septic shock conditions (narrow), Anaphylactic/anaphylactoid shock conditions (narrow), Hypoglycaemic and neurogenic shock conditions (narrow), Conditions associated with central nervous system haemorrhages and cerebrovascular accidents (narrow), Hypotonic-hyporesponsive episode (broad), Chronic kidney disease (narrow), Hypersensitivity (narrow), Drug reaction with eosinophilia and systemic symptoms syndrome (broad), Infective pneumonia (broad), Sepsis (narrow), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
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: 03/07/21 SARS-CoV-2 (COVID-19) by NAA, Micro Not Detected; 03/25/21 SARS-CoV-2 (COVID-19) by NAA, Micro Not Detected; 05/08/21 SARS-CoV-2 (COVID-19) by NAA, Micro DETECTED
CDC Split Type:

Write-up: Pt. w/ "multiple medical problems including failed renal transplant x 2 with ESRD on HD, SAH, HTN, OSA 2018, afib and acute DVT in Dec 20 on eliquis, orthostatic hypotension, adrenal insuff (secondary), recent pseudomonas sepsis who presented to facility on 3/25/2021 with infection of L leg wound. Was sent from surgeon''s office when noted to have increased pain, increased drainage from wound. Left lower extremity wound with exposed muscle and tendons."... "initially transferred to a different department on 4/23/21 but transferred off on 5/8/21 due to cough and fevers to 104.4 attributed to new diagnosis of COVID-19." Pt. died on 08/24/21 in hospice, after D/C from ICU due to MRSA bacteremia,

ESRD, Arrest, Shock, Septic Emboli to the brain.

VAERS ID: 1691329 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: Unknown
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-11

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

Administered by: Other **Purchased by:** ?
Symptoms: SARS-CoV-2 test, Suspected COVID-19, Vaccination failure
SMQs:, Lack of efficacy/effect (narrow), Infective pneumonia (broad), Opportunistic infections (broad), COVID-19 (narrow)

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 2021-08-01
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data: Test Name: COVID-19 virus test; Result Unstructured Data: Covid-19
CDC Split Type: USJNJFOC20210913396

Write-up: SUSPECTED COVID-19 INFECTION; SUSPECTED CLINICAL VACCINATION FAILURE; This spontaneous report received from a media via a company representative concerned an elderly patient (over 80 years) of an unspecified sex. The patient''s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported) dose was not reported, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced suspected covid-19 infection and suspected clinical vaccination failure. On an unspecified date (over the past week of 03-SEP-2021) in AUG-2021, the patient died from suspected covid-19 infection. It was unknown if an autopsy was performed or not. Laboratory data (dates unspecified) included: COVID-19 virus test (NR: not provided) Covid-19. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of suspected covid-19 infection on an unspecified date, and the

outcome of suspected clinical vaccination failure was not reported. This report was serious (Death, and Other Medically Important Condition). This report was associated with product quality complaint: 90000192684.; Sender"s Comments: V0: 20210913396- Covid-19 vaccine ad26.cov2.s-Suspected covid-19 infection. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210913396- Covid-19 vaccine ad26.cov2.s-Suspected clinical vaccination failure. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: SPECIAL SITUATIONS; Reported Cause(s) of Death: SUSPECTED COVID-19 INFECTION

VAERS ID: 1691338 (history)
Form: Version 2.0
Age:
Sex: Unknown
Location: New York
Vaccinated: 0000-00-00
Onset: 0000-00-00
Submitted: 0000-00-00
Entered: 2021-09-11

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

Administered by: Other **Purchased by:** ?
Symptoms: Death
SMQs:

Life Threatening? No
Birth Defect? No
Died? Yes
Date died: 0000-00-00
Permanent Disability? No
Recovered? No
Office Visit? No
ER Visit? No
ER or Doctor Visit? No
Hospitalized? No
Previous Vaccinations:
Other Medications:
Current Illness:
Preexisting Conditions:
Allergies:
Diagnostic Lab Data:
CDC Split Type: USJNJFOC20210914084

Write-up: DEATH DUE TO THE VACCINE; This spontaneous report received from a consumer concerned a 32 year old of unspecified sex. The patient"s height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic

vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died due to the Johnson and Johnson vaccine. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender"s Comments: V0: 20210914084-Covid-19 vaccine ad26.cov2.s-Death due to the vaccine. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

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