The Vaccine Adverse Event Reporting System (VAERS) Results

VAERS ID	Adverse Event Description	Lab Data	Current Illness	Adverse Events After Prior Vaccinations	Medications At Time Of Vaccination	History/Allergies
1400303-1	After the second dose she had immediate pain at the injection site and over the next 24 hours she developed: A fever of 101.4 severe abdominal pain and chest pain that made her feel like her heart was being pulled out painful electric shocks down her neck and spine that made her walk hunched over numbness and swelling in the arm she got the shot pain in her fingers and toes that turned white and were ice cold to the touch Over the next 2.5 months her abdominal, muscle and nerve pain became unbearable plus she developed new symptoms: Fatigue gastroparesis, nausea and vomiting Eventually she couldn?t even swallow food or liquids without immediately spitting it up. An itchy rash on her arms peeling skin on her feet Her menstrual cycle lasted a month with large clumps of blood She had unexplained painful cysts vision problems headaches erratic blood pressure and heart rate memory loss, mixing up words and brain fog Dizziness, fainting and then nonepileptic seizures that we suspect were from lyrica verbal and motor tics loss of feeling from the waist down, muscle weakness, abnormal gait and eventually she wasn?t able to walk at all urinary retention From the day she got her 2nd dose to today we took her to the ER nine (9) times and she was admitted to the hospital a total of 3 times totalling 2 months. The last time she was	Multiple blood tests, spine MRI, Upper GI, Endoscopy, x-ray and ultrasound of abdomen. Can provide test results, too much to type in here. Additional blood work done today and brain MRI/MRV scheduled for 6/22.	none	No prior vaccinations for this event.	Vyvanse 50mg	dermatographia,none before the vaccine, tape allergy after the vaccine

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	admitted to the hospital she could not walk, was unable to feel or move below her waist, threw up anything she tried to eat or drink, had tachycardia and her blood sugar was at 47. Once she got an NG tube and was stable they transferred her to Inpatient Rehabilitation and she was just discharged on June 1st. Today she is able to walk with a walker and take care of herself but she still has little to no feeling below her waist. She still has an NG tube for nutrition and continues to have GI and urinary retention problems.					

Note: Submitting a report to VAERS does not mean that healthcare personnel or the vaccine caused or contributed to the adverse event (possible side effect).

Notes:

Caveats: VAERS accepts reports of adverse events and reactions that occur following vaccination. Healthcare providers, vaccine manufacturers, and the public can submit reports to VAERS. While very important in monitoring vaccine safety, VAERS reports alone cannot be used to determine if a vaccine caused or contributed to an adverse event or illness. The reports may contain information that is incomplete, inaccurate, coincidental, or unverifiable. Most reports to VAERS are voluntary, which means they are subject to biases. This creates specific limitations on how the data can be used scientifically. Data from VAERS reports should always be interpreted with these limitations in mind.

The strengths of VAERS are that it is national in scope and can quickly provide an early warning of a safety problem with a vaccine. As part of CDC and FDA's multi-system approach to post-licensure vaccine safety monitoring, VAERS is designed to rapidly detect unusual or unexpected patterns of adverse events, also known as "safety signals." If a safety signal is found in VAERS, further studies can be done in safety systems such as the CDC's Vaccine Safety Datalink (VSD) or the Clinical Immunization Safety Assessment (CISA) project. These systems do not have the same limitations as VAERS, and can better assess health risks and possible connections between adverse events and a vaccine.

Key considerations and limitations of VAERS data:

- Vaccine providers are encouraged to report any clinically significant health problem following vaccination to VAERS, whether or not they believe the vaccine was the cause.
- Reports may include incomplete, inaccurate, coincidental and unverified information.
- The number of reports alone cannot be interpreted or used to reach conclusions about the existence, severity, frequency, or rates of problems associated with vaccines.
- VAERS data are limited to vaccine adverse event reports received between 1990 and the most recent date for which data are available.
- VAERS data do not represent all known safety information for a vaccine and should be interpreted in the context of other scientific information.

Some items may have more than 1 occurrence in any single event report, such as Symptoms, Vaccine Products, Manufacturers, and Event Categories. If data are grouped by any of these items, then the number in the Events Reported column may exceed the total number of unique events. If percentages are shown, then the associated percentage of total unique event reports will exceed 100% in such cases. For example, the number of Symptoms mentioned is likely to exceed the number of events reported, because many reports include more than 1 Symptom. When more then 1 Symptom occurs in a single report, then the percentage of Symptoms to unique events is more than 100%. More information. (/wonder/help/vaers.html#Suppress)

Data contains VAERS reports processed as of 07/02/2021. The VAERS data in WONDER are updated weekly, yet the VAERS system receives continuous updates including revisions and new reports for preceding time periods. Duplicate event reports and/or reports determined to be false are removed from VAERS. More information. (/wonder/help/vaers.html#Reporting)

For more information on how many persons have been vaccinated in the US for COVID19 to date, see https://covid.cdc.gov/covid-data-tracker/#vaccinations/ (https://covid.cdc.gov/covid-data-tracker/#vaccinations/)

Help: See The Vaccine Adverse Event Reporting System (VAERS) Documentation (/wonder/help/vaers.html) for more information.

Query Date: Jul 13, 2021 10:19:25 PM

Suggested Citation:

United States Department of Health and Human Services (DHHS), Public Health Service (PHS), Centers for Disease Control (CDC) / Food and Drug Administration (FDA), Vaccine Adverse Event Reporting System (VAERS) 1990 - 07/02/2021, CDC WONDER On-line Database. Accessed at http://wonder.cdc.gov/vaers.html on Jul 13, 2021 10:19:25 PM

Query Criteria:

Age:	6-17 years
State / Territory:	Ohio
Vaccine Lot:	220395
Vaccine Manufacturer:	PFIZER\BIONTECH
Vaccine Products:	COVID19 VACCINE (COVID19)
VAERS ID:	All
Group By:	VAERS ID
Show Totals:	False
Show Zero Values:	False