

# More Than 1.3 Million Adverse Events Following COVID Vaccines Reported to VAERS, CDC Data Show

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*VAERS data released Friday by the Centers for Disease Control and Prevention show 1,329,135 reports of adverse events from all age groups following COVID-19 vaccines, including 29,273 deaths and 241,910 serious injuries between Dec. 14, 2020, and July 1, 2022.*

The Centers for Disease Control and Prevention (CDC) today released new data showing a total of [1,329,135 reports of adverse events](#) following [COVID-19](#) vaccines were submitted between Dec. 14, 2020, and July 1, 2022, to the Vaccine Adverse Event Reporting System (VAERS). That’s an increase of 14,541 adverse events over the previous week.

VAERS is the primary government-funded system for reporting adverse vaccine reactions in the U.S.

The data included a total of [29,273 reports of deaths](#) — an increase of 111 over the previous week — and [241,910 serious injuries](#), including deaths, during the same time period — up 684 compared with the previous week.

Of the 29,273 reported deaths, [18,937 cases](#) are attributed to Pfizer’s COVID-19 vaccine, [7,724 cases](#) to Moderna and [2,545 cases](#) to Johnson & Johnson (J&J).

Excluding “[foreign reports](#)” to VAERS, [839,450 adverse events](#), including [13,547 deaths](#) and [85,321 serious injuries](#), were reported in the U.S. between Dec. 14, 2020, and July 1, 2022.

[Foreign reports](#) are reports foreign subsidiaries send to U.S. vaccine manufacturers. Under U.S. Food and Drug Administration (FDA) regulations, if a manufacturer is notified of a foreign case report that describes an event that is both serious and does not appear on the product’s labeling, the manufacturer is required to submit the report to VAERS.

Of the 13,547 U.S. [deaths reported](#) as of July 1, 15% occurred within 24 hours of

vaccination, 19% occurred within 48 hours of vaccination and 58% occurred in people who experienced an [onset of symptoms](#) within 48 hours of being vaccinated.

In the U.S., 596 million COVID-19 vaccine doses had been administered as of June 29, [including](#) 352 million doses of Pfizer, 225 million doses of Moderna and 19 million doses of Johnson & Johnson (J&J).



National Vaccine Information Center  
Your Health. Your Family. Your Choice.

## Search Results

From the 7/1/2022 release of VAERS data:

### Found 1,329,135 cases where Vaccine is COVID19

[Government Disclaimer on use of this data](#)

Table

↓	↑ ↓	↑ ↓
Event Outcome	Count	Percent
Death	29,273	2.2%
Permanent Disability	54,693	4.11%
Office Visit	197,404	14.85%
Emergency Room	121	0.01%
Emergency Doctor/Room	131,295	9.88%
Hospitalized	165,313	12.44%
Hospitalized, Prolonged	403	0.03%
Recovered	347,951	26.18%
Birth Defect	1,120	0.08%
Life Threatening	32,600	2.45%
Not Serious	614,857	46.26%
<b>TOTAL</b>	† 1,575,030	† 118.5%

† Because some cases have multiple vaccinations and symptoms, a single case can account for multiple entries in this table. This is why the Total Count is greater than 1,329,135 (the number of cases found), and the Total Percent is greater than 100.

Every Friday, [VAERS](#) publishes vaccine injury reports received as of a specified date. Reports submitted to VAERS require further investigation before a causal relationship can be confirmed.

Historically, VAERS has been shown to report only [1% of actual vaccine adverse events](#).

#### U.S. VAERS data from Dec. 14, 2020, to July 1, 2022, for 6-month-olds to 5-year-olds show:

- 1,941 [adverse events](#), including 64 cases [rated as serious](#) and 3 [reported deaths](#).
- 4 [reports](#) of myocarditis and pericarditis (heart inflammation).  
The CDC uses a [narrowed case definition](#) of “myocarditis,” which [excludes cases](#) of cardiac arrest, [ischemic strokes](#) and deaths due to heart problems that occur before one has the chance to go to the emergency department.
- 13 [reports](#) of blood clotting disorders.

#### U.S. VAERS data from Dec. 14, 2020, to July 1, 2022, for 5- to 11-year-olds show:

- 11,638 [adverse events](#), including 300 [rated as serious](#) and 7 [reported deaths](#).

- [23 reports](#) of myocarditis and pericarditis.
- [46 reports](#) of blood clotting disorders.

#### **U.S. VAERS data from Dec. 14, 2020, to July 1, 2022, for 12- to 17-year-olds show:**

- [32,543 adverse events](#), including [1,843 rated as serious](#) and [44 reported deaths](#).
- [62 reports](#) of anaphylaxis among 12- to 17-year-olds where the reaction was life-threatening, required treatment or resulted in death — with 97% of cases attributed to [Pfizer’s vaccine](#).
- [655 reports](#) of myocarditis and pericarditis with [643 cases](#) attributed to Pfizer’s vaccine.  
There was one less case reported attributed to Pfizer’s vaccine since the previous week.
- [166 reports](#) of blood clotting disorders with all cases attributed to Pfizer. VAERS reported 167 cases of blood clotting disorders in the 12- to 17-year-old age group last week.
- [20 cases](#) of postural orthostatic tachycardia syndrome (POTS) with [all cases](#) attributed to Pfizer’s vaccine.

#### **U.S. VAERS data from Dec. 14, 2020, to July 1, 2022, for all age groups combined, show:**

- 20% of deaths were related to cardiac disorders.
- 54% of those who died were male, 41% were female and the remaining death reports did not include the gender of the deceased.
- The [average age](#) of death was 73.
- As of July 1, [5,629 pregnant women](#) reported adverse events related to COVID-19 vaccines, including [1,761 reports of miscarriage or premature birth](#).
- Of the [3,623 cases of Bell’s Palsy](#) reported, 51% were attributed to [Pfizer](#) vaccinations, 40% to [Moderna](#) and 8% to [J&J](#).
- [895 reports of Guillain-Barré syndrome](#), with 42% of cases [attributed to Pfizer](#), 30% to [Moderna](#) and 27% to [J&J](#).
- [2,285 reports](#) of anaphylaxis where the reaction was life-threatening, required treatment or resulted in death.
- [1,733 reports](#) of myocardial infarction.
- [14,171 reports](#) of blood-clotting disorders in the U.S. Of those, [6,334 reports](#) were attributed to Pfizer, [5,088 reports](#) to Moderna and [2,713 reports](#) to J&J.
- [4,251 cases](#) of myocarditis and pericarditis with [2,606 cases](#) attributed to Pfizer, [1,443 cases](#) to Moderna and [187 cases](#) to J&J.
- [13 cases](#) of Creutzfeldt-Jakob disease with [7 cases](#) attributed Pfizer, [5 cases](#) to Moderna and [1 case](#) to J&J.
- [268 cases](#) of POTS with [165 cases](#) attributed to Pfizer, [84 cases](#) to Moderna and [17 cases](#) to J&J.

### **Uruguay halts COVID vaccine for kids under 13, judge demands government officials turn over Pfizer contracts**

Uruguay [suspended COVID-19 vaccines](#) for children under 13 after a judge on Thursday issued an injunction halting vaccinations in that age group until government officials turn over contracts with vaccine manufacturers.

Uruguayan government officials and Pfizer were [ordered on Wednesday](#) to appear in court after a judge gave them 48 hours to present detailed information on Pfizer's COVID-19 vaccine while the court considered an injunction request to halt COVID-19.

The government said a confidentiality clause in the contract prevents it from producing the documents, [The Washington Post reported](#).

**Judge Alejandro Recarey** of the Administrative Litigation Tribunal used his [inquisitorial powers](#) to demand the Uruguayan Ministry of Public Health, State Health Services Administration and the President's Office submit all information regarding the contracts for the purchase of COVID-19 vaccines, including contractual information related to any clauses of civil indemnity or criminal impunity of the suppliers in the event of adverse effects.

The judge also [posed a series of questions](#) to government officials and Pfizer regarding the chemical composition, efficacy and safety of COVID-19 vaccines, and required Pfizer to state whether it has "admitted, in any area, internal or external to it and its partners, the verification of adverse effects" of its COVID-19 vaccines in children.

## **Paul Offit says 'fix was in' before FDA panel voted to reformulate boosters**

In a June 6 [interview](#) with ZDoggMD, vaccine expert **Dr. Paul Offit** said the FDA's vaccine advisory panel's recent meeting on whether to modify COVID-19 boosters was unusual and he felt the panel was led to "vote yes" to reformulate boosters without critical data.

Offit, director of the Vaccine Education Center and professor of pediatrics in the Division of Infectious Diseases at Children's Hospital of Philadelphia, said:

"I've seen nothing like this. I guess the thing that's most upsetting to me is normally when you get something from the FDA when we have these meetings, you usually get it a few days before you meet. You usually get a couple of hundred pages.

"Here on the other hand, normally you get the EUA [Emergency Use Authorization] submission from the company, which is 85 to 100 pages long, and then you get the FDA's review of all those data. It's a very thorough review. Not here though. Here, it was 22 pages from the FDA, which included a half-page on Pfizer's data and a half-page on Moderna's data."

"You could get that from the [press release](#)," Offit said, adding he didn't see the benefits of reformulated boosters and was surprised that out of 21 voting members, [19 voted "yes."](#)

Offit also said it was "unusual" that someone from the World Health Organization presented their opinion during the meeting that a modified booster shot was a good idea and the FDA presented their opinion before the advisory committee rendered its advice.

Offit said he believed reformulating COVID-19 boosters was "something that was desired by the Biden administration," who announced the day after the meeting they had [purchased at least 105 million doses](#) from Pfizer with up to 300 million doses."

[Children's Health Defense](#) asks anyone who has experienced an adverse reaction, to any vaccine, to file a report following [these three steps](#).

## **76 doctors urge UK government officials not to authorize COVID vaccines for small children**

In a [letter](#) to the Medical and Healthcare products Regulatory Agency (MHRA) and other U.K. government officials, 76 doctors explain why the recent U.S. Food and Drug Administration's (FDA) decision authorizing COVID vaccinations for infants and young children must not happen in the U.K.

The doctors [urged the MHRA](#) to consider "very carefully" the move to vaccinate ever younger children against SARS-CoV-2, despite the gradual but significant reducing virulence of successive variants, the increasing evidence of rapidly waning vaccine efficacy, increasing concerns over long-term vaccine harms and the knowledge that the vast majority of the younger age group have already been exposed to SARS-CoV-2 repeatedly and have demonstrably effective immunity.

The doctors said using the risk-benefit analysis that supported the rollout of mRNA vaccines to the elderly and vulnerable in 2021 is totally inappropriate for small children in 2022, and strongly challenged the addition of COVID-19 vaccination into the [routine child immunization program](#) despite no demonstrated clinical need, known and unknown risks and the fact that these vaccines still have only conditional marketing authorization.

## **60,000 unvaccinated guard and reserve soldiers cut from training and pay**

About 60,000 Army National Guard members and Army Reserve soldiers who [refused to comply](#) with a Department of Defense COVID-19 vaccine mandate are no longer allowed to participate in their military duties and were cut off from some of their pay and benefits, Army officials announced July 1.

Of the more than 40,000 members of the Guard who remain unvaccinated, 14,000 have said they do not intend to ever receive a COVID-19 vaccine, Guard officials [told CBS News](#). Approximately 22,000 Reserve soldiers have refused to get vaccinated.

Soldiers [will be allowed](#) to come on duty and earn their pay if it's for the purpose of getting vaccinated or to take part in separation procedures. If the soldiers continue to refuse to get vaccinated, the consequences could be even more severe.

To date, only six Guard soldiers across all states and territories have received medical exemptions out of 53 who submitted requests, according to Army data. No Reserve soldiers have received a medical exemption.

No Guard or Reserve soldiers have been approved for a religious exemption despite nearly 3,000 requests.

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