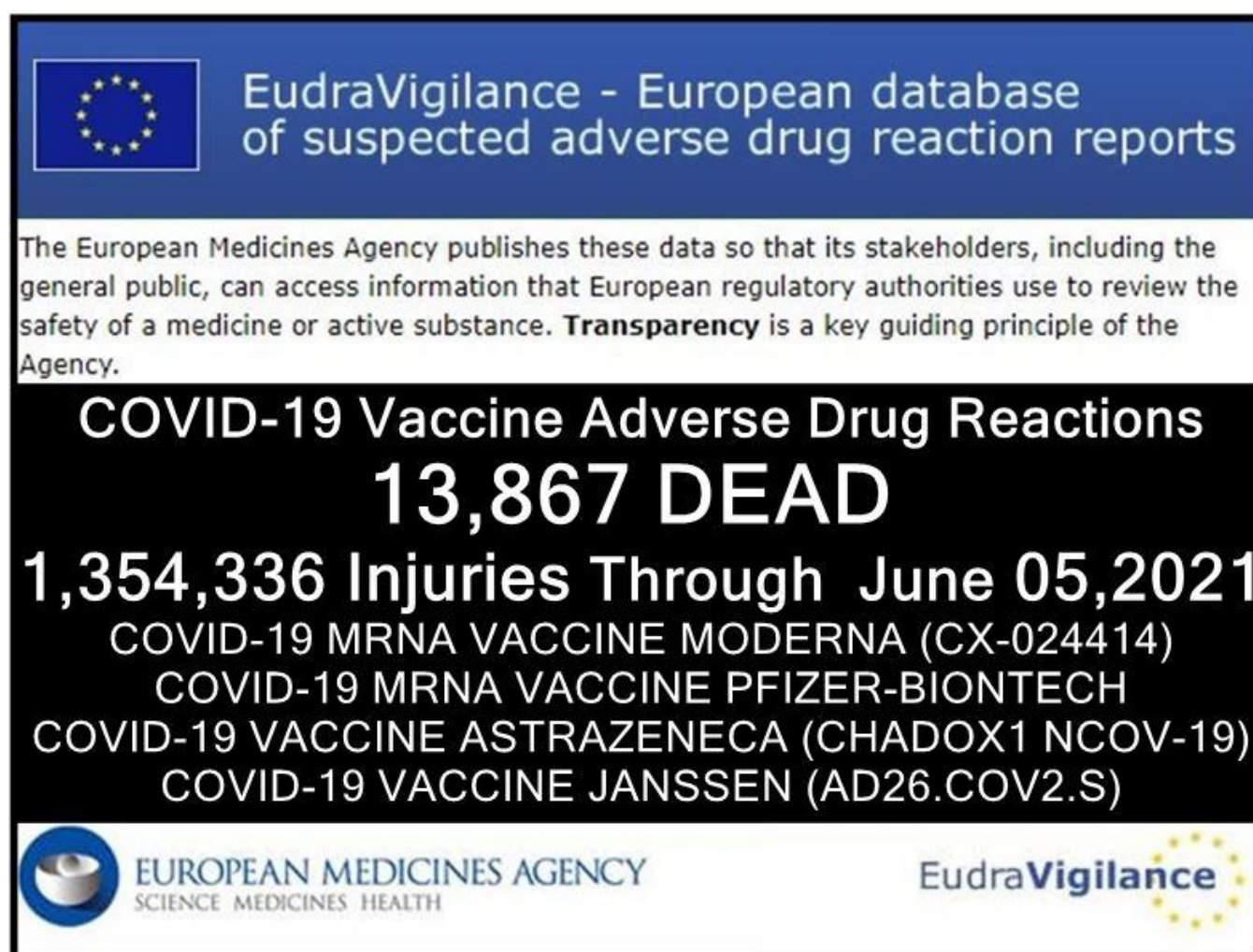


Worldwide Genocide Continues: 13,867 DEAD and 1,354,336 Injuries in European Database of Adverse Drug Reactions for COVID- 19 Shots

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The banner features the European Union flag logo on the left. The text is as follows:


**EudraVigilance - European database
of suspected adverse drug reaction reports**

The European Medicines Agency publishes these data so that its stakeholders, including the general public, can access information that European regulatory authorities use to review the safety of a medicine or active substance. **Transparency** is a key guiding principle of the Agency.

**COVID-19 Vaccine Adverse Drug Reactions
13,867 DEAD
1,354,336 Injuries Through June 05, 2021**

COVID-19 MRNA VACCINE MODERNA (CX-024414)
COVID-19 MRNA VACCINE PFIZER-BIONTECH
COVID-19 VACCINE ASTRAZENECA (CHADOX1 NCOV-19)
COVID-19 VACCINE JANSSEN (AD26.COV2.S)

 **EUROPEAN MEDICINES AGENCY**
SCIENCE MEDICINES HEALTH

EudraVigilance 

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The European database of suspected drug reaction reports is which also

tracks reports of injuries and deaths following the experimental COVID-19 "vaccines."

Here is what states about their database:

*This website was launched by the in 2012 **to provide public access to reports of suspected side effects** (also known as suspected adverse drug reactions). These reports are submitted electronically to by national medicines regulatory authorities and by pharmaceutical companies that hold marketing authorisations (licences) for the medicines.*

EudraVigilance is a system designed for collecting reports of suspected side effects. These reports are used for evaluating the benefits and risks of medicines during their development and monitoring their safety following their authorisation in the European Economic Area (EEA). EudraVigilance has been in use since December 2001.

This website was launched to comply with the , which was developed to improve public health by supporting the monitoring of the safety of medicines and to increase transparency for stakeholders, including the general public.

The first approved the EudraVigilance Access Policy in December 2010. A revision was adopted by the Board in December 2015 based on the 2010 pharmacovigilance legislation. The policy aims to provide stakeholders such as national medicines regulatory authorities in the EEA, the European Commission, healthcare professionals, patients and consumers, as well as the pharmaceutical industry and research organisations, with access to reports on suspected side effects.

Transparency is a key guiding principle of the Agency, and is pivotal to building trust and confidence in the regulatory process. By increasing

transparency, the Agency is better able to address the growing need among stakeholders, including the general public, for access to information. (.)

Their report through June 5, 2021 lists **13,867 deaths and 1,354,336 injuries** following injections of four experimental COVID-19 shots:

From the total of injuries recorded, there are 683,688 **serious** injuries which equals over 50%.

*“**Seriousness** provides information on the suspected undesirable effect; it can be classified as ‘serious’ if it corresponds to a medical occurrence that results in **death**, is life-threatening, requires inpatient hospitalisation, results in another medically important condition, or prolongation of existing hospitalisation, results in persistent or significant disability or incapacity, or is a congenital anomaly/birth defect.”*

A *Health Impact News* subscriber in Europe ran the reports for each of the four COVID-19 shots we are including here. This subscriber has volunteered to do this, and it is a lot of work to tabulate each reaction with injuries and fatalities, since there is no place on the system we have found that tabulates all the results.

Since we have started publishing this, others from Europe have also calculated the numbers and confirmed the totals.*

Here is the summary data through June 5, 2021.

Total reactions for the experimental mRNA vaccine **Tozinameran** (code **BNT162b2, Comirnaty**) from **BioNTech/Pfizer: 6,732 deaths** and **502,162 injuries** to 05/06/2021

Total reactions for the experimental mRNA vaccine **mRNA-1273(CX-024414)** from **Moderna: 3,821 deaths** and **101,767 injuries** to

05/06/2021

Total reactions for the experimental vaccine AZD1222/VAXZEVRIA (CHADOX1 NCOV-19) from Oxford/ AstraZeneca: 2,848 deaths and 724,457 injuries to 05/06/2021

Total reactions for the experimental COVID-19 vaccine JANSSEN (AD26.COVS.S) from Johnson & Johnson: 466 deaths and 25,950 injuries to 05/06/2021

Last Update: Jun 05 2021	Reported Cases	Fatalities	% fatalities to cases	All Multiple Symptoms	Serious injuries	% serious to ALL
Astrazeneca	264 549	2 848	1,08%	724 457	404 748	55,87%
Pfizer-BioNTech	212 053	6 732	3,17%	502 162	210 964	42,01%
Moderna	40 712	3 821	9,39%	101 767	58 174	57,16%
Jannsen	8 593	466	5,42%	25 950	9 802	37,77%
Total:	525 907	13 867	2,64%	1 354 336	683 688	50,48%

*These totals are estimates based on reports submitted to . Totals may be much higher based on percentage of adverse reactions that are reported. Some of these reports may also be reported to the individual country's adverse reaction databases, such as the U.S. VAERS database, and the UK Yellow Card system. The fatalities are grouped by symptoms, and some fatalities may have resulted from multiple symptoms.

See Also: