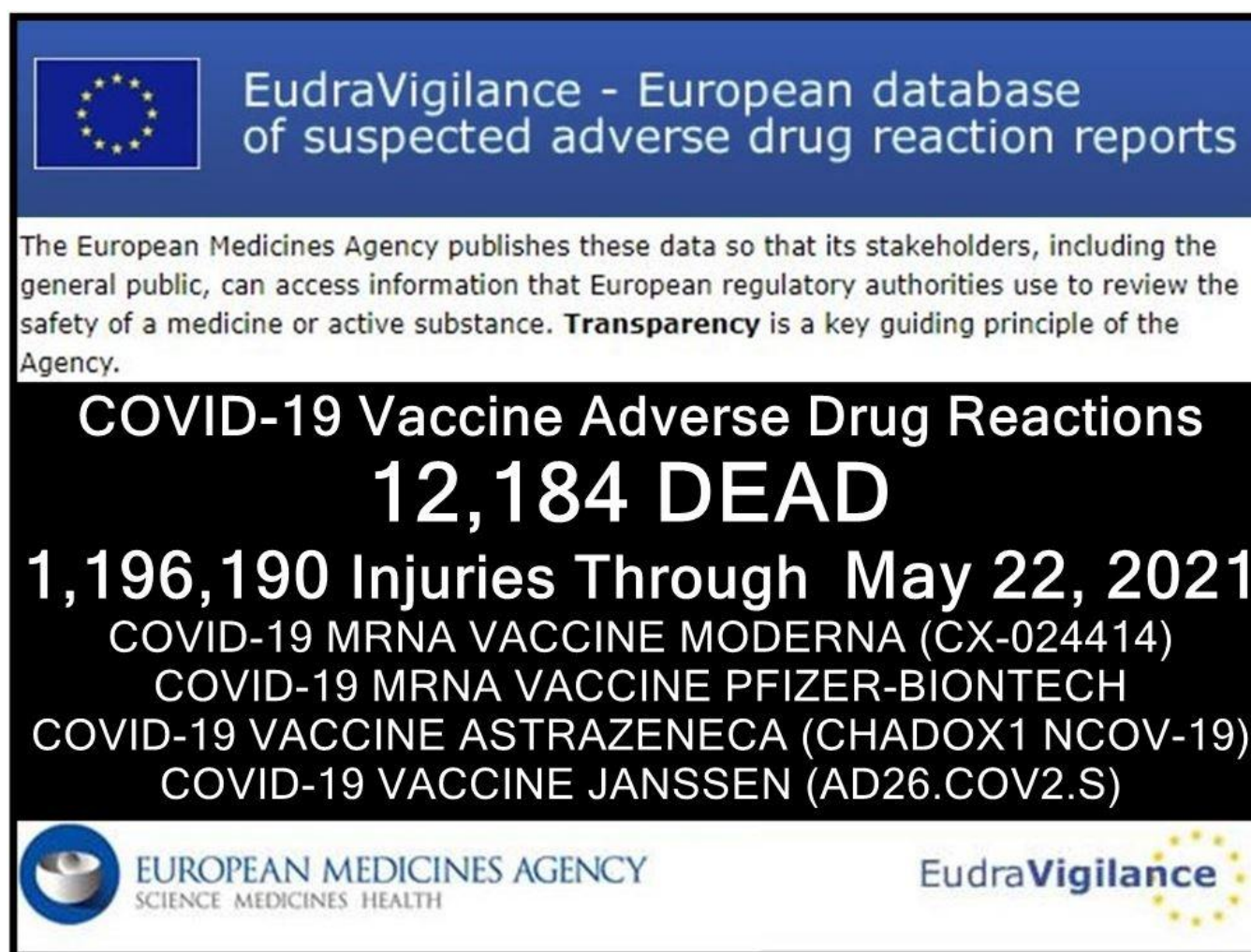


12,184 DEAD 1,196,190 Injuries: European Database of Adverse Drug Reactions for COVID-19 "Vaccines"

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The screenshot shows the EudraVigilance website header with the European Union flag and the text "EudraVigilance - European database of suspected adverse drug reaction reports". Below this, a paragraph states: "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." The main content area has a black background with white text that reads: "COVID-19 Vaccine Adverse Drug Reactions", "12,184 DEAD", "1,196,190 Injuries Through May 22, 2021", and lists four vaccine types: "COVID-19 MRNA VACCINE MODERNA (CX-024414)", "COVID-19 MRNA VACCINE PFIZER-BIONTECH", "COVID-19 VACCINE ASTRAZENECA (CHADOX1 NCOV-19)", and "COVID-19 VACCINE JANSSEN (AD26.COV2.S)". The footer includes the European Medicines Agency logo and the EudraVigilance logo.

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
12,184 DEAD
1,196,190 Injuries Through May 22, 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

by **Brian Shilhavy**
Editor, Health Impact News

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 EMA 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 EU Directive, 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 EMA 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 May 22, 2021 lists **12,184 deaths and 1,196,190**

injuries following injections of four experimental COVID-19 shots:

From the total of injuries recorded, there are 604,744 **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 May 22, 2021.

Total reactions for the experimental mRNA vaccine **Tozinameran** (code **BNT162b2, Comirnaty**) from **BioNTech/_Pfizer**: **5,961 deaths** and **452,779 injuries** to 22/05/2021

Total reactions for the experimental mRNA vaccine **mRNA-1273(CX-024414)** from **Moderna**: **3,365 deaths** and **72,596 injuries** to 22/05/2021

Total reactions for the experimental vaccine **AZD1222/VAXZEVRIA (CHADOX1 NCOV-19)** from **Oxford/ AstraZeneca**: **2,489 deaths** and **655,534 injuries** to 22/05/2021

Total reactions for the experimental COVID-19 vaccine JANSSEN (AD26.COV2.S) from Johnson & Johnson: 369 deaths and 15,281 injuries to 22/05/2021

Last Update: May 22 2021	Reported Cases	Fatalities	All Multiple Symptoms	Serious injuries	
Astrazeneca	237 648	2 489	655 534	372 019	56,75%
Pfizer-BioNTech	191 215	5 961	452 779	186 308	41,15%
Moderna	29 616	3 365	72 596	38 704	53,31%
Janssen	4 997	369	15 281	7 713	50,47%
Total:	463 476	12 184	1 196 190	604 744	50,56%

*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: