

# HOUSE BILL 1189

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By: **Delegates M. Fisher, Arentz, Boteler, Krebs, Mautz, Morgan, Rose, Szeliga, and Thiam**

Introduced and read first time: February 11, 2022

Assigned to: Health and Government Operations

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## A BILL ENTITLED

1 AN ACT concerning

2 **Public Health – Vaccines Administered Under Emergency Use Authorization –**  
3 **Reporting of Adverse Events**

4 FOR the purpose of requiring a health care provider to report to the national Vaccine  
5 Adverse Event Reporting System an adverse event occurring after the  
6 administration of a vaccine, including a COVID–19 vaccine or booster, administered  
7 under an emergency use authorization; and generally relating to reporting adverse  
8 events related to vaccines administered under emergency use authorization.

9 BY adding to

10 Article – Health – General

11 Section 18–405

12 Annotated Code of Maryland

13 (2019 Replacement Volume and 2021 Supplement)

14 SECTION 1. BE IT ENACTED BY THE GENERAL ASSEMBLY OF MARYLAND,  
15 That the Laws of Maryland read as follows:

16 **Article – Health – General**

17 **18–405.**

18 **(A) (1) IN THIS SECTION THE FOLLOWING WORDS HAVE THE MEANINGS**  
19 **INDICATED.**

20 **(2) “ADVERSE EVENT” MEANS ANY ILLNESS, DISABILITY,**  
21 **INCAPACITY, HOSPITALIZATION, DEATH, OR IMPAIRMENT OF MENTAL, EMOTIONAL,**  
22 **BEHAVIORAL, OR PHYSICAL FUNCTIONING OR DEVELOPMENT, THE FIRST**  
23 **MANIFESTATION OF WHICH APPEARS AFTER THE DATE OF ADMINISTRATION OF A**

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EXPLANATION: CAPITALS INDICATE MATTER ADDED TO EXISTING LAW.

[Brackets] indicate matter deleted from existing law.



1 VACCINE, INCLUDING A COVID-19 VACCINE OR BOOSTER.

2 (3) "HEALTH CARE PROVIDER" MEANS AN INDIVIDUAL WHO IS  
3 LICENSED, CERTIFIED, OR OTHERWISE AUTHORIZED UNDER THE HEALTH  
4 OCCUPATIONS ARTICLE TO PROVIDE HEALTH CARE SERVICES.

5 (4) "VAERS" MEANS THE NATIONAL VACCINE ADVERSE EVENT  
6 REPORTING SYSTEM ADMINISTERED BY THE U.S. CENTERS FOR DISEASE CONTROL  
7 AND PREVENTION AND THE U.S. FOOD AND DRUG ADMINISTRATION.

8 (B) A HEALTH CARE PROVIDER SHALL PROMPTLY REPORT TO VAERS ANY  
9 ADVERSE EVENT OBSERVED IN A PATIENT AFTER THE ADMINISTRATION OF A  
10 VACCINE, INCLUDING A COVID-19 VACCINE OR BOOSTER, ADMINISTERED UNDER  
11 AN EMERGENCY USE AUTHORIZATION.

12 SECTION 2. AND BE IT FURTHER ENACTED, That this Act shall take effect  
13 October 1, 2022.