

PRESS RELEASE

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10-31-2023



Defending The Republic's FOIA Case Against The FDA Uncovers New Data on Adverse Events

Defendant the Republic (DTR) has just obtained another set of documents from the Food and Drug Administration (FDA) relating to the agency's approval of Moderna's COVID-19 vaccine "Spikevax." DTR received these Moderna files as a result of its ongoing Freedom of Information Act (FOIA) lawsuit against the FDA. They are available for download below.

The new records include a listing of adverse events for thousands of clinical participants who received doses of Moderna's COVID-19 vaccine. The adverse events, many of which started mere days after vaccination, include: elevated blood pressure, swollen lymph nodes, high fevers and fatigue, rashes, and vertigo.

These are just some of the adverse reactions that clinical trial participants had to Moderna's COVID-19 vaccine. In July of 2023, DTR **released** the initial set of Moderna documents – the first significant release of data from Moderna's COVID-19 clinical trials – which revealed serious adverse events following the company's vaccine, including death and neurological disorders.

Also released are: summaries of the duration of days that passed from the date of injection (for both the first and second shots) and the onset of adverse reactions; Moderna responses to FDA comments concerning vaccine effectiveness; and other FDA comments to Moderna concerning the clinical results.

Before the end of the year, DTR anticipates the FDA, in compliance with its FOIA obligations, will produce thousands more pages of Moderna records. We will summarize and release those records to the public once available.

Please **support** our efforts to expose government corruption, promote government accountability and transparency, fight COVID-19 mandates on behalf of military servicemembers, combat censorship, and to restore integrity in our elections.

[Moderna Oct 2023 Document Production](#)

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