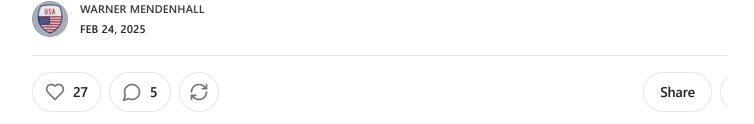
Did Covid "vaccines" violate environmental safety law?

A petition was filed 1-21-2025 with the FDA by Julian Gillespie, Kevin McKernan, Dr. Dav Speicher, Dr. Jessica Rose, and L. Maria Gutschi.



I had not examined environmental assessment criteria in any litigation. I look forwato the FDA response in the new administration. This <u>citizen petition</u> was filed Janu 21, 2025 with the FDA requesting administrative action regarding Pfizer and Moderna's COVID-19 mRNA vaccines to:

- 1. Revoke or suspend the Biologics License Applications (BLAs) for Pfizer (Comirnaty) and Moderna (Spikevax) COVID-19 vaccines.
- 2. Investigate regulatory processes and decisions.
- 3. Issue public guidance on risks of synthetic DNA contamination.
- 4. Mandate independent testing of existing stocks.
- 5. Establish testing for DNA integration into human cells.

Primary Concerns:

Wrongful categorical exclusions from Environmental Assessments (EAs)

- 1. Excessive synthetic DNA contamination, with multiple studies showing levels 470 times above regulatory safety thresholds.
- 2. Presence of SV40 promoter and enhancer sequences.

3. Potential risks including genomic integration and cancer.

Legal Arguments:

- 1. The products should have been classified as gene therapies.
- 2. The FDA wrongfully granted categorical exclusions from environmental assessments.
- 3. The BLA approvals were legally void from the beginning due to these failures.

They argue these issues represent significant public health risks requiring immedia regulatory action and investigation.

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