Gmail **Christine, an unincorporated woman <cmssyc@gmail.com>**

CDC confesses: no scientific basis for "HPV" narrative... no virus, no contagion, no genome... just bogus tests

Christine, an unincorporated woman <cmssyc@gmail.com> Wed, Oct 16, 2024 at 4:48 PM
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resending to include an omitted section:

Notice to agent is notice to principal; notice to principal is notice to agent...in your public and private capacities

August 29, 2024:
The “experts” at the **U.S. Centers for Disease Control and Prevention** and the **Agency for Toxic Substances and Disease Registry** were officially challenged via a freedom of information order (**pgs 1/2**) to provide or cite any studies in their possession, custody or control ***authored by anyone, anywhere, ever***:

1. that **scientifically prove/provide evidence of the existence** of any alleged "HPV" aka "Human Papillomavirus" (showing that the alleged particles exist and cause the illness/symptoms that they are alleged to cause), or

2. that even describe someone **finding and purifying particles** alleged to be "HPV" directly from the bodily fluid/tissue/excrement of so-called "hosts", or

3. wherein the **purported "genome" of the alleged "HPV" was found intact** in the bodily fluid, tissue or excrement of any supposedly infected “host” (as opposed to fabricated in silico, aka a computer model), or

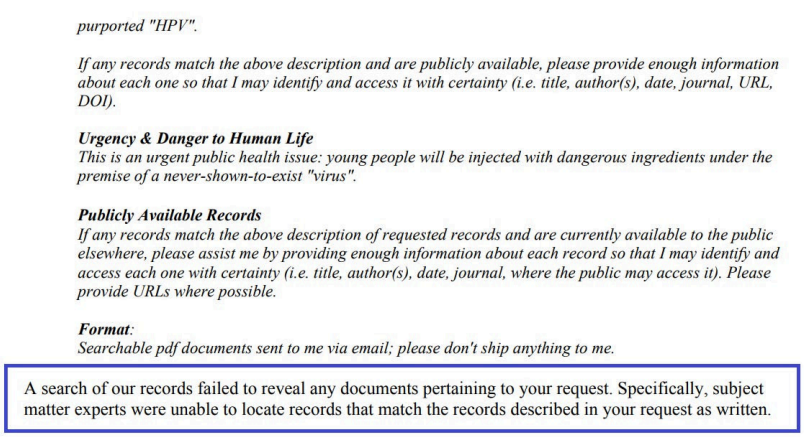
4. that **scientifically demonstrate contagion** of the illness / symptoms that are allegedly caused by purported "HPV".

I included a reminder that scientific evidence requires use of the scientific method to test falsifiable hypotheses through valid, rigorous, repeatable controlled experiments.

And as usual I asked that if records matching my request were held by the institutions but were already publicly available, I be given citations.

October 16, 2024:
Roger Andoh acting as FOIA Officer in the Office of the Chief Operating Officer responded (**pgs 7, 8**, #24-01612-FOIA):

> “A search of our records failed to reveal any documents pertaining to your request. Specifically, subject matter experts were unable to locate records that match the records described in your request as written.”



purported "HPV".

If any records match the above description and are publicly available, please provide enough information about each one so that I may identify and access it with certainty (i.e. title, author(s), date, journal, URL, DOI).

Urgency & Danger to Human Life
This is an urgent public health issue: young people will be injected with dangerous ingredients under the premise of a never-shown-to-exist "virus".

Publicly Available Records
If any records match the above description of requested records and are currently available to the public elsewhere, please assist me by providing enough information about each record so that I may identify and access each one with certainty (i.e. title, author(s), date, journal, where the public may access it). Please provide URLs where possible.

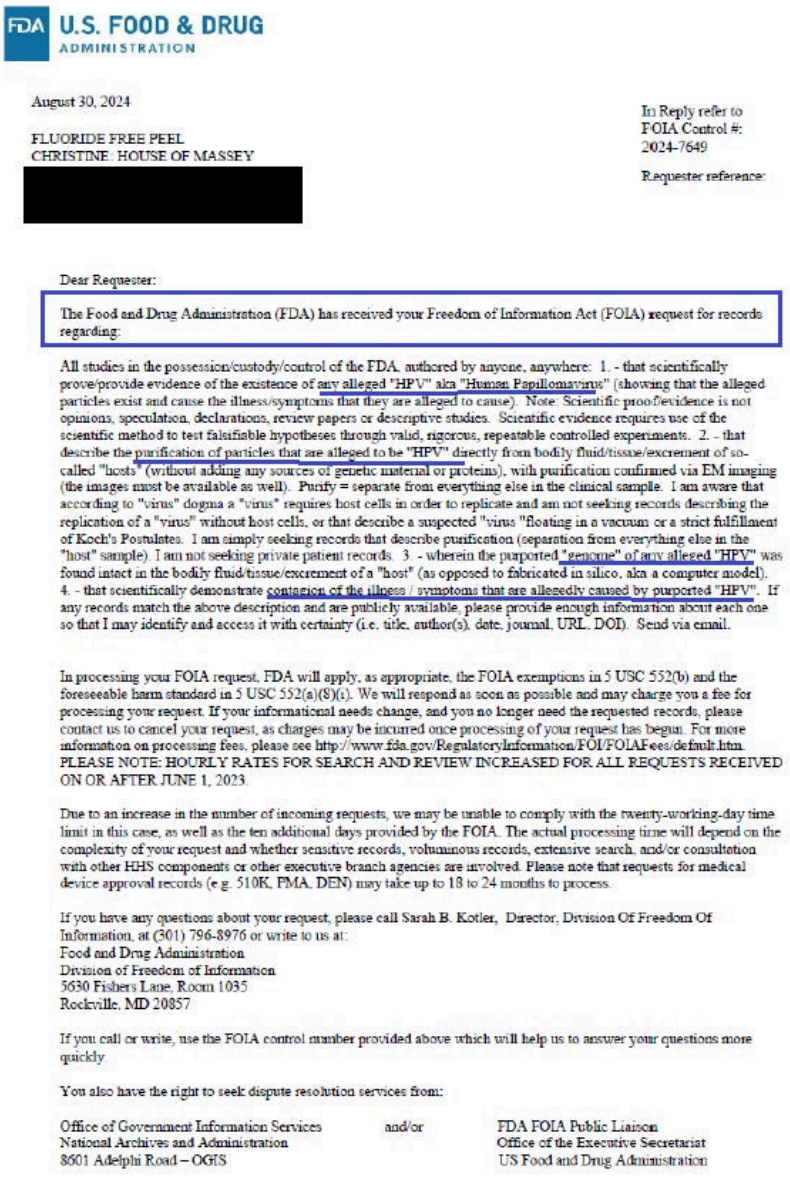
Format:
Searchable pdf documents sent to me via email; please don't ship anything to me.

A search of our records failed to reveal any documents pertaining to your request. Specifically, subject matter experts were unable to locate records that match the records described in your request as written.

And now we await the same confession from the FDA in response to FOIA order #2024-7649, of which Sarah Kolter acting as Director acknowledged receipt on August 30, 2024.

Note that Sarah has already officially confessed that the FDA has no such records for the imaginary “monkeypox virus” (FDA FOIA **2024-7353**) or the imaginary “avian influenza virus” (FDA FOIA **2024-6486**) and she was not able to cite any for the imaginary “SARS-COV-2” either (FDA FOIA **2024-7837**)**…**

…because virology has always been **pseudoscience**, no virus has **ever** been shown to exist, contagion is “public health” **mythology** and literally **hundreds** of earlier FOI responses from 40 different countries on dozens of alleged “viruses” also yielded no valid scientific evidence (see further below for more links).



FDA U.S. FOOD & DRUG ADMINISTRATION

August 30, 2024

FLUORIDE FREE PEEL
CHRISTINE HOUSE OF MASSEY
[redacted]

In Reply refer to
FOIA Control #:
2024-7649

Requester reference:

Dear Requester:

The Food and Drug Administration (FDA) has received your Freedom of Information Act (FOIA) request for records regarding:

All studies in the possession/custody/control of the FDA, authored by anyone, anywhere: 1. - that scientifically prove/provide evidence of the existence of any alleged "HPV" aka "Human Papillomavirus" (showing that the alleged particles exist and cause the illness/symptoms that they are alleged to cause). Note: Scientific proof/evidence is not opinions, speculation, declarations, review papers or descriptive studies. Scientific evidence requires use of the scientific method to test falsifiable hypotheses through valid, rigorous, repeatable controlled experiments. 2. - that describe the purification of particles that are alleged to be "HPV" directly from bodily fluid/tissue/excrement of so-called "hosts" (without adding any sources of genetic material or proteins), with purification confirmed via EM imaging (the images must be available as well). Purify = separate from everything else in the clinical sample. I am aware that according to "virus" dogma a "virus" requires host cells in order to replicate and am not seeking records describing the replication of a "virus" without host cells, or that describe a suspected "virus "floating in a vacuum or a strict fulfillment of Koch's Postulates. I am simply seeking records that describe purification (separation from everything else in the "host" sample). I am not seeking private patient records. 3. - wherein the purported "genome" of any alleged "HPV" was found intact in the bodily fluid/tissue/excrement of a "host" (as opposed to fabricated in silico, aka a computer model). 4. - that scientifically demonstrate contagion of the illness / symptoms that are allegedly caused by purported "HPV". If any records match the above description and are publicly available, please provide enough information about each one so that I may identify and access it with certainty (i.e. title, author(s), date, journal, URL, DOI). Send via email.

In processing your FOIA request, FDA will apply, as appropriate, the FOIA exemptions in 5 USC 552(b) and the foreseeable harm standard in 5 USC 552(a)(8)(i). We will respond as soon as possible and may charge you a fee for processing your request. If your informational needs change, and you no longer need the requested records, please contact us to cancel your request, as charges may be incurred once processing of your request has begun. For more information on processing fees, please see http://www.fda.gov/RegulatoryInformation/FOI/FOIAFees/default.htm. PLEASE NOTE: HOURLY RATES FOR SEARCH AND REVIEW INCREASED FOR ALL REQUESTS RECEIVED ON OR AFTER JUNE 1, 2023.

Due to an increase in the number of incoming requests, we may be unable to comply with the twenty-working-day time limit in this case, as well as the ten additional days provided by the FOIA. The actual processing time will depend on the complexity of your request and whether sensitive records, voluminous records, extensive search, and/or consultation with other HHS components or other executive branch agencies are involved. Please note that requests for medical device approval records (e.g. 510K, PMA, DEN) may take up to 18 to 24 months to process.

If you have any questions about your request, please call Sarah B. Kotler, Director, Division Of Freedom Of Information, at (301) 796-8976 or write to us at:
Food and Drug Administration
Division of Freedom of Information
5630 Fishers Lane, Room 1035
Rockville, MD 20857

If you call or write, use the FOIA control number provided above which will help us to answer your questions more quickly.

You also have the right to seek dispute resolution services from:

Office of Government Information Services
National Archives and Administration
8601 Adelphi Road – OGIS

and/or

FDA FOIA Public Liaison
Office of the Executive Secretariat
US Food and Drug Administration

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