

FOIA Request Confirmation

Confirmation Number: FDA24106575

Requester:

General							
Description of Requester: Educational institution or noncommercial scientific institution operated primarily for scholarly or scientific research							
Max Amount Willing to Pay: \$40							
Organization							
Organization Name: Fluc	oride Free Peel						
Primary Phone:		Other Phone:		Email:	cmssyc@gmail.com		
Mailing Address Billing Address							
Address 1			Address 1:				
Address 2			Address 2:				
City			City:				
Province CA			Province:				
Postal Code CA			Postal Code:				
Country			Country:				
Details							
Requester Name: Christine: House of Massey							
Reque	ster File #:	Request Letter		Request Letter:			
Requested D	Date From: 01/01/1945			Requested Date To: 08/28/2024			
Subject of Request:		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					
Cubject of Hodgest.	"HPV" aka "Hum	"HPV" aka "Human Papillomavirus" (showing that the alleged particles exist and cause the illness/symptoms that they are alleged to cause). Note: Scientific					
Waiver of Fees							
Justification: Public interest, public health issue of great importance.							
Expedited Processing							
Reason: D	Reason: Danger to human life						
Justification: P	Justification: Public interest, urgent public health issue: young people will be injected with dangerous ingredients under the premise of a never-shown-to-exist "virus".						

Print Create Another Request Close

Within 10 business days of the submission of your online request, you will receive by electronic mail an FOIA Control Number. If you need to communicate with FDA regarding your request, please refer to this Control Number. Requests received after 4:00 P.M. E.S.T. will be considered to have been received on the following business day.



FOI Request Received

noreply@fda.gov <noreply@fda.gov>
To: cmssyc@gmail.com

Thu, Aug 29, 2024 at 6:37 PM

*** This is an automated message. Please do not reply to this email. ***

Reference: FDA24106575

Dear Requester,

This is to confirm that you submitted a request for record(s) from the Food and Drug Administration pursuant to the Freedom of Information Act.

FOIA staff will review your request to determine whether it has sufficient information to be processed; if so, you will receive another email as a formal acknowledgement of your request, with a control number for your request. If your request is not sufficiently described, or if there are any other deficiencies with your submission, FOIA staff will contact you via telephone or email.



FDA Receipt of FOI Request Control # 2024-7649

FDA_FOI@fda.gov <FDA_FOI@fda.gov>

Fri, Aug 30, 2024 at 7:44 AM

To: cmssyc@gmail.com Cc: FDAFOIA@fda.hhs.gov

Control number: 2024-7649

Please find the attached acknowledgement regarding your FOIA request.

Note: Do NOT reply directly to this E-mail





August 30, 2024

FLUORIDE FREE PEEL CHRISTINE: HOUSE OF MASSEY 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 socalled "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 College Park, MD 20740-6001 Telephone:202-741-5770 Toll-Free: 1-877-684-6448

Email:ogis@nara.gov Fax: 202-741-5769 5630 Fishers Lane, Room 1050 Rockville, MD 20857 Email: FDAFOIA@fda.hhs.gov

Sincerely,

SARAH KOTLER Director 11/30/24, 1:19 PM Gmail - 2024-10300



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

2024-10300

Christine, an unincorporated woman <cmssyc@gmail.com> To: "Kotler, Sarah" <Sarah.Kotler@fda.hhs.gov>

Sat, Nov 30, 2024 at 1:18 PM

Thank you Sarah.

I'm still waiting for your response re HPV, see attached.

Christine

[Quoted text hidden]



2024 08 30 emailed Acknowledgement Letter.PDF 62K



FDA Receipt of FOI Request Control # 2024-7649

Christine, an unincorporated woman <cmssyc@gmail.com>
To: FDA_FOI@fda.gov, FDAFOIA@fda.hhs.gov, "Kotler, Sarah" <sarah.kotler@fda.hhs.gov>

Mon, Jan 6, 2025 at 10:39 PM

Good evening,

I've been waiting over 4 months for the response. Kindly send it in a letter, signed with Sarah's proper name.

Christine
[Quoted text hidden]



FDA Receipt of FOI Request Control # 2024-7649

Kotler, Sarah <Sarah.Kotler@fda.hhs.gov>

Tue, Jan 7, 2025 at 7:40 AM

To: "Christine, an unincorporated woman" <cmssyc@gmail.com>, "FDA_FOI@fda.gov" <FDA_FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

You have acknowledged receipt of our response. I am not clear on what you mean by "Sarah's proper name." My signature information is below and includes my full proper name.

Sarah B. Kotler, J.D.

Director, Division of Headquarters Freedom of Information

Office of Management and Enterprise Services

Office of the Commissioner

US Food & Drug Administration

301-796-8976

Sarah.Kotler@fda.hhs.gov

















From: Christine, an unincorporated woman <cmssyc@gmail.com>

Sent: Monday, January 6, 2025 10:39 PM



FDA Receipt of FOI Request Control # 2024-7649

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

Tue, Jan 7, 2025 at 2:01 PM

To: "Kotler, Sarah" <Sarah.Kotler@fda.hhs.gov>

Cc: "FDA FOI@fda.gov" <FDA FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

Correction:

I acknowledged your unsigned acknowledgement letter that stated "We will respond as soon as possible....", not your legal confession that "the experts" at FDA have zero scientific evidence of the imaginary HPV or even the imaginary HPV "genome" and no scientific evidence that the symptoms purportedly caused by the imaginary virus are contagious because you have not supplied said confession yet.

In your Poxviridae confession (published here and attached) you "digitally signed" as "Sarah B. Kotler -S".

You did the same in your "SARS-COV-2" confession, attached and published here.

You did the same in your "monkeypox virus" confession, attached and published here (and you ignored all of my questions).

You provided your "avian influenza virus" confession as an email without even a "digital signature", attached and published here.

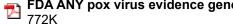
I will publish an article about this request in the next week, with or without your final response.

The longer you stall, the worse it looks and the longer you might spend in jail for fraud, culpable homicide = murder, etc.

Christine

[Quoted text hidden]

4 attachments



FDA ANY pox virus evidence genome contagion PACKAGE redacted.pdf





FDA avian influenza virus no records 2024 07 PACKAGE redacted.pdf 1042K



FDA Receipt of FOI Request Control # 2024-7649

Kotler, Sarah <Sarah.Kotler@fda.hhs.gov>

Tue, Jan 7, 2025 at 2:16 PM

To: "Christine, an unincorporated woman" <cmssyc@gmail.com>

Cc: "FDA_FOI@fda.gov" <FDA_FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

Pursuant to 45 CFR 5.28(a), "we will send you a response informing you of our release determination, including whether any responsive records were located, how much responsive material was located, whether the records are being released in full or withheld in full or in part, any fees you must pay for processing of the request, and your right to seek assistance from the appropriate FOIA Public Liaison." We have done that. Your request is closed. Are you saying you did not receive the response? If that is the case, I can send you another one.

Thanks,



FDA Receipt of FOI Request Control # 2024-7649

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

Tue, Jan 7, 2025 at 2:32 PM

To: "Kotler, Sarah" <Sarah.Kotler@fda.hhs.gov>

Cc: "FDA_FOI@fda.gov" <FDA_FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

No response to this FOIA order ever appeared in my inbox.



FDA Receipt of FOI Request Control # 2024-7649

Kotler, Sarah <Sarah.Kotler@fda.hhs.gov>

Tue, Jan 7, 2025 at 2:34 PM

To: "Christine, an unincorporated woman" <cmssyc@gmail.com>

Cc: "FDA_FOI@fda.gov" <FDA_FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

Ok, now I understand what you are asking. We will send a new response.



FDA Receipt of FOI Request Control # 2024-7649

Kotler, Sarah <Sarah.Kotler@fda.hhs.gov>

Tue, Jan 7, 2025 at 2:43 PM

To: "Christine, an unincorporated woman" <cmssyc@gmail.com>

Cc: "FDA FOI@fda.gov" <FDA FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

By email to: cmssyc@gmail.com

FDA FOIA 2024-7649

Dear Requester,

I am writing in response to your request for:

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.

The FDA does not regulate or treat viruses. The FDA is responsible for protecting the public health by ensuring the safety, efficacy, and security of human and veterinary drugs, biological products, and medical devices; and by ensuring the safety of our nation's food supply, cosmetics, and products that emit radiation. For general information on HVP, please see: HPV (human papillomavirus) | FDA.

In accordance with 45 CFR § 5.61 and 21 CFR § 20.41(b)(5), you have the right to appeal this determination. Your appeal should clearly identify the agency determination that is being appealed. It would be helpful if you provide specific reasons explaining why you believe the agency's adverse determination should be reconsidered. By filing an appeal, you preserve your rights under FOIA and give the agency a chance to review and reconsider your request and the agency's decision. Your appeal must be mailed within 90 days from the date of this response, to: Director, Office of the Executive Secretariat, U.S. Food & Drug Administration, 5630 Fishers Lane, Room 1050, Rockville, MD 20857, or emailed within 90 days from the date of this response to FDAFOIA@fda.hhs.gov. Please clearly mark both the envelope and your letter "FDA Freedom of Information Act Appeal." Items arriving or delivered after 5 p.m. Eastern Time will be deemed received on the next workday.

If you would like to discuss our response <u>before</u> filing an appeal to attempt to resolve your dispute without going through the appeals process, please contact me at 301-796-8976 or <u>Sarah.Kotler@fda.hhs.gov</u>.

Please contact me if you have questions about your request. You may also contact the FDA FOIA Public Liaison for assistance at: Office of the Executive Secretariat, US Food & Drug Administration, 5630 Fishers Lane, Room 1050, Rockville, MD 20857, E-mail: FDAFOIA@fda.hhs.gov.

If you are unable to resolve your FOIA dispute through our FOIA Public Liaison, the Office of Government Information Services (OGIS), the Federal FOIA Ombudsman's office, offers mediation services to help resolve disputes between FOIA

requesters and Federal agencies. The contact information for OGIS is as follows: Office of Government Information Services, National Archives and Records Administration, 8601 Adelphi Road–OGIS, College Park, MD 20740-6001; telephone at 202-741-5770; toll free at 1-877-684-6448; or facsimile at 202-741-5769; e-mail at ogis@nara.gov.

Sincerely,



FDA Receipt of FOI Request Control # 2024-7649

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

Tue, Jan 7, 2025 at 2:59 PM

To: "Kotler, Sarah" <Sarah.Kotler@fda.hhs.gov>

Cc: "FDA FOI@fda.gov" <FDA FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

Thank you but I don't see any information in that email regarding "release determination, including whether any responsive records were located, how much responsive material was located, whether the records are being released in full or withheld in full or in part". It does not address my FOIA order.

And these comments are irrelevant, false and misleading so kindly remove them:

"The FDA does not regulate or treat viruses." - No one can regulate or treat alleged viruses because they have never been shown to exist. Both the literature and literally hundreds of FOI responses from 40 different countries confirm this (as shown at the links below and in my notarized affidavit, attached).

"The FDA is responsible for protecting the public health by ensuring the safety, efficacy, and security of human and veterinary drugs, biological products, and medical devices; and by ensuring the safety of our nation's food supply, cosmetics, and products that emit radiation." - The FDA rubber stamps poisons like quackcines and is complicit in pseudoscience and deadly fraud. The nation's food supply is largely contaminated with filth of various sorts.

"For general information on HVP, please see: HPV (human papillomavirus) | FDA." - There is no information about any alleged virus, only mis- and dis-information about imaginary viruses. No virus has ever been shown to exist.

The whole point of my FOIA orders has been to give "the experts" the chance to prove that no-virus people are wrong. "The experts" at FDA have failed in every instance, just like "the experts" at every other institution that was challenged.

Christine

[Quoted text hidden]

7

2024 08 12 virus FOIs affidavit NOTARTIZED REDACTED.pdf 904K



FDA Receipt of FOI Request Control # 2024-7649

Kotler, Sarah <Sarah.Kotler@fda.hhs.gov>

Tue, Jan 7, 2025 at 3:19 PM

To: "Christine, an unincorporated woman" <cmssyc@gmail.com>

Cc: "FDA_FOI@fda.gov" <FDA_FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

We consider your request misdirected. It is not something that FDA would have records about. You may want to consider submitting to CDC. You are also welcome to appeal, contact the FOIA Public Liaison, or contact OGIS.



FDA Receipt of FOI Request Control # 2024-7649

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

Tue, Jan 7, 2025 at 4:35 PM

To: "Kotler, Sarah" < Sarah.Kotler@fda.hhs.gov>

Cc: "FDA FOI@fda.gov" <FDA FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

So now you admit that you willfully chose not to provide a response (because there is no valid scientific evidence for you to provide or cite, and "the experts" and others, including yourself, at FDA are complicit in a massive deadly hoax), apparently in violation of 45 CFR 5.28(a). You could have just admitted that from the outset, instead of pretending that you had already sent a response, and then sending me a no-response fake-response.

There is nothing "misdirected" about challenging fake-experts and fake-authorities to provide or cite valid scientific evidence to back up their deadly, fraud-based hoaxes.

As explained to you in the past and in my affidavit, the CDC has already been formally challenged on dozens of alleged viruses, and they also failed to provide or cite even a shred of valid scientific evidence for any of them, or even a "genome" or contagion of symptoms.

I will not waste my time on an appeal since your last email already made it clear for everyone to see: **you know perfectly well** that no one at the FDA can provide or cite any valid evidence of any alleged virus or "viral genome" or "viral" anything or contagion. The fraudulent rubber-stamping of impossible-to-validate tests, deadly quackcines, etc. at the FDA is just part of the murderous theatre.

Life in prison!

Christine



Tue, Jan 7, 2025 at 4:51 PM

FDA Receipt of FOI Request Control # 2024-7649

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

To: "Kotler, Sarah" <Sarah.Kotler@fda.hhs.gov>

Cc: "FDA_FOI@fda.gov" <FDA_FOI@fda.gov>, FDA FOIA <FDAFOIA@fda.hhs.gov>

p.s. Still waiting for your explanation of your strange signatures. [Quoted text hidden]