U.S. Food and Drug Administration

FOIA Request Confirmation

Confirmation Number: FDA24108874

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:	**Fluoride Free Peel**				
Primary Phone:	[redacted]	Other Phone:		Email:	**cmssyc@gmail.com**

Mailing Address

Address 1:	[redacted]
Address 2:	
City:	
Province CA:	
Postal Code CA:	
Country:	

Billing Address

Address 1:	[redacted]
Address 2:	
City:	
Province:	
Posta Code:	
Country:	

Details

Requester Name:	**Christine Massey**		
Requester File #:		Request Letter:	
Requested Date From:	**01/01/1900**	Requested Date To:	**11/19/2024**
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**		

Waiver of Fees

Justification:	**Matter of public health interest.**

Expedited Processing

Reason:	
Justification:	

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.

If your informational needs change, and you need to cancel your request, please contact the Division of Freedom of Information by telephone, mail, or fax. Please include your control number in the correspondence. For contact information, please see FDA's FOIA page.

FDA U.S. FOOD & DRUG ADMINISTRATION

November 21, 2024

FLUORIDE FREE PEEL
CHRISTINE MASSEY
[redacted]

In Reply refer to
FOIA Control #:
2024-10300

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 virus of the "Poxviridae family" (showing at minimum that the alleged particles with a specific "genome" and proteins 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. - or, that even describe purification of particles claimed to be any of said alleged viruses directly from bodily fluid/tissue/excrement of so-called "hosts" (without adding any sources of genetic material or proteins) with purification confirmed via EM images (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. - or, wherein the purported "genome" of any of these alleged viruses was found intact in the bodily fluid/tissue/excrement of a so-called "host" (as opposed to fabricated/assembled in silico, aka a computer model) 4. - or, that scientifically demonstrate contagion of any of the illnesses allegedly caused by any of these alleged viruses. Send via email. I do not agree to this order or "personal information" being shared with 3rd parties including those that provide FOIA services, without my explicit written consent.

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

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

2024-10300

Kotler, Sarah <Sarah.Kotler@fda.hhs.gov> Fri, Nov 29, 2024 at 10:07 AM
To: "Christine, an unincorporated woman" <cmssyc@gmail.com>

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 virus of the "Poxviridae family" (showing at minimum that the alleged particles with a specific "genome" and proteins 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. - or, that even describe purification of particles claimed to be any of said alleged viruses directly from bodily fluid/tissue/excrement of so-called "hosts" (without adding any sources of genetic material or proteins) with purification confirmed via EM images (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. - or, wherein the purported "genome" of any of these alleged viruses was found intact in the bodily fluid/tissue/excrement of a so-called "host" (as opposed to fabricated/assembled in silico, aka a computer model) 4. - or, that scientifically demonstrate contagion of any of the illnesses allegedly caused by any of these alleged viruses. Send via email. I do not agree to this order or "personal information" being shared with 3rd parties including those that provide FOIA services, without my explicit written consent.

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. Therefore, we have no responsive records. If you are interested, the NIH has information about this topic at: Poxviruses - Medical Microbiology - NCBI Bookshelf.

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 before filing an appeal to attempt to resolve your dispute without going through the appeals process, please contact me at 301-796-8976 or Sarah.Kotler@fda.hhs.gov.

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,

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

OMES

November 29.pdf
205K

U.S. FOOD & DRUG ADMINISTRATION

November 29, 2024

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Sincerely,

Sarah B. Kotler -S

Digitally signed by Sarah B. Kotler -S
Date: 2024.11.29 10:06:18 -05'00'

Sarah Kotler

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

2024-10300

Christine, an unincorporated woman <cmssyc@gmail.com> Sat, Nov 30, 2024 at 2:06 PM
To: "Kotler, Sarah" <Sarah.Kotler@fda.hhs.gov>

P.S.
Sarah, why is your letter "Digitally signed" by "Sarah B. Kotler -S"? Please provide a letter with your proper name.
[Quoted text hidden]