Exhibit (C):
August 9, 2021 Email and Acknowledgement Letter

Complaint for Injunctive Relief,
Reed Smith LLP v. Food & Drug Administration
Civil Action No. 21-2387

Sylora, Andrew J.

From: FDA_FOI@fda.gov
Sent: Monday, August 9, 2021 6:47 AM
To: Pontikes, Rachael G.
Cc: FDAFOIA@fda.hhs.gov
Subject: FDA Receipt of FOI Request Control # 2021-5130
Attachments: Acknowledgement Letter.PDF

EXTERNAL E-MAIL - From FDA_FOI@fda.gov

Control number: 2021-5130

Please find the attached acknowledgement regarding your FOIA request.

Note: Do NOT reply directly to this E-mail

U.S. FOOD & DRUG ADMINISTRATION

August 09, 2021

REED SMITH LLP
RACHAEL G. PONTIKES
10 South Wacker Drive
Chicago IL 60606 USA

In Reply refer to
FOIA Control #:
2021-5130

Requester reference:

Dear Requester:

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

We request the following records: (1) all communications* between FDA (including, but not limited to, FDA's Center for Drug Evaluation and Research ("CDER")) and the Pharmacy Compounding Advisory Committee from January 1, 2014, to the present regarding the following subjects: (b) nicotinamide adenine dinucleotide; or ETC

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.

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 Rochelle A. Coleman, Information Technician, at (301) 796-8982 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
College Park, MD 20740-6001
Telephone:202-741-5770
Toll-Free: 1-877-684-6448
Email:ogis@nara.gov
Fax: 202-741-5769

and/or

FDA FOIA Public Liaison
Office of the Executive Secretariat
US Food Administration
5630 Fishers Lane, Room 1050
Email: FDAFOIA@fda.hhs.gov

Sincerely,

SARAH KOTLER
Director