Comments that you submit in response to this notice are a matter of public record. Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

Abstract: The BIE provides standards for the appropriate use of lands and facilities by third parties under 25 CFR part 48. These standards address the following: the execution of lease agreements; the establishment and administration of mechanisms for the acceptance of consideration for the use and benefit of a Bureau-operated school; the assurance of ethical conduct; and monitoring the amount and terms of consideration received, the manner in which the consideration is used, and any results achieved by such use.

Title of Collection: Use of Bureau-Operated Schools by Third Parties.

OMB Control Number: 1076–0187.

Form Number: None.

Type of Review: Extension of a currently approved collection.

Respondents/Affected Public: Individuals and private sector.

Total Estimated Number of Annual Respondents: 17.

Total Estimated Number of Annual Responses: 24.

Estimated Completion Time per Response: One to three hours.

Total Estimated Number of Annual Burden Hours: 68 hours.

Respondent's Obligation: Required to Obtain a Benefit.

Frequency of Collection: Annually.

Total Estimated Annual Nonhour Burden Cost: $0.

An agency may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

The authority for this action is the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*).

Steven Mullen,

Information Collection Clearance Officer, Office of Regulatory Affairs and Collaborative Action—Indian Affairs.

[FR Doc. 2024–13403 Filed 6–17–24; 8:45 am]

BILLING CODE 4337–15–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1313]

Certain Botulinum Toxin Products and Processes for Manufacturing or Relating to Same; Notice of Request for Submissions on the Public Interest

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that on June 10, 2024, the presiding administrative law judge (''ALJ'') issued an Initial Determination on Violation of Section 337. On June 10, 2024, the ALJ also issued a Recommended Determination on remedy and bonding should a violation be found in the above-captioned investigation. The Commission is soliciting submissions on public interest issues raised by the recommended relief should the Commission find a violation. This notice is soliciting comments from the public and interested government agencies only.

FOR FURTHER INFORMATION CONTACT: Michael Liberman, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–3115. Copies of non-confidential documents filed in connection with this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.* For help accessing EDIS, please email *EDIS3Help@usitc.gov.* General information concerning the Commission may also be obtained by accessing its internet server at *https://www.usitc.gov.* Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: Section 337 of the Tariff Act of 1930 provides that, if the Commission finds a violation, it shall exclude the articles concerned from the United States unless, after considering the effect of such exclusion upon the public health and welfare, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, and United States consumers, it finds that such articles should not be excluded from entry. (19 U.S.C. 1337(d)(1)). A similar provision applies to cease and desist orders. (19 U.S.C. 1337(f)(1)).

The Commission is soliciting submissions on public interest issues raised by the recommended relief should the Commission find a violation, specifically: (1) a limited exclusion order directed to certain botulinum toxin products and processes for manufacturing or relating to same imported, sold for importation, and/or sold after importation by respondents Hugel, Inc. of Seoul, Republic of Korea; Hugel America, Inc. of Irvine, California; and Croma Pharma GmbH of Leobendorf, Austria; and (2) a cease and desist order directed to certain botulinum toxin products and processes for manufacturing or relating to same imported, sold for importation, and/or sold after importation by respondent Hugel America, Inc. of Irvine, California. Parties are to file public interest submissions pursuant to 19 CFR 210.50(a)(4).

The Commission is interested in further development of the record on the public interest in this investigation. Accordingly, members of the public and interested government agencies are invited to file submissions of no more than five (5) pages, inclusive of attachments, concerning the public interest in light of the ALJ's Recommended Determination on Remedy and Bonding issued in this investigation on June 10, 2024. Comments should address whether issuance of the recommended remedial orders in this investigation, should the Commission find a violation, would affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

(i) explain how the articles potentially subject to the recommended remedial orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the recommended orders;

(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;

(iv) indicate whether complainant, complainant's licensees, and/or third-party suppliers have the capacity to replace the volume of articles potentially subject to the recommended orders within a commercially reasonable time; and

(v) explain how the recommended orders would impact consumers in the United States.

Written submissions must be filed no later than by close of business on July 12, 2024.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. The Commission's paper filing requirements in 19 CFR 210.4(f) are currently waived. 85 FR 15798 (Mar. 19, 2020). Submissions should refer to the investigation number (''Inv. No. 337–TA–1313'') in a prominent place on the cover page and/or the first page. (*See Handbook for Electronic Filing Procedures, https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf.*). Persons with questions regarding filing should contact the Secretary (202–205–2000).

Any person desiring to submit a document to the Commission in confidence must request confidential treatment by marking each document with a header indicating that the document contains confidential information. This marking will be deemed to satisfy the request procedure set forth in Rules 201.6(b) and 210.5(e)(2) (19 CFR 201.6(b) & 210.5(e)(2)). Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. Any non-party wishing to submit comments containing confidential information must serve those comments on the parties to the investigation pursuant to the applicable Administrative Protective Order. A redacted non-confidential version of the document must also be filed simultaneously with any confidential filing and must be served in accordance with Commission Rule 210.4(f)(7)(ii)(A) (19 CFR 210.4(f)(7)(ii)(A)). All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. appendix 3; or (ii) by U.S. Government employees and contract personnel, solely for cybersecurity purposes. All contract personnel will sign appropriate nondisclosure agreements. All nonconfidential written submissions will be available for public inspection on EDIS.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: June 12, 2024.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2024–13333 Filed 6–17–24; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1377]

Certain Products Containing Tirzepatide and Products Purporting To Contain Tirzepatide; Notice of a Commission Determination Not To Review an Initial Determination Granting a Motion To Amend the Complaint and Notice of Investigation

AGENCY: International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission (''Commission'') has determined not to review an initial determination (''ID'') (Order No. 16) of the presiding administrative law judge (''ALJ'') granting a motion to amend the complaint and notice of investigation to name an additional respondent.

FOR FURTHER INFORMATION CONTACT: Edward S. Jou, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–3316. Copies of non-confidential documents filed in connection with this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.* For help accessing EDIS, please email *EDIS3Help@usitc.gov.* General information concerning the Commission may also be obtained by accessing its internet server at *https://www.usitc.gov.* Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal, telephone (202) 205–1810.

SUPPLEMENTARY INFORMATION: The Commission instituted this investigation on November 27, 2023, based upon a complaint filed on behalf of Eli Lilly and Company (''Eli Lilly'') of Indianapolis, Indiana. 88 FR 82914–15 (Nov. 27, 2023). The complaint, as supplemented, alleges violations of section 337 based upon the importation into the United States and the sale of certain products containing tirzepatide or purporting to contain tirzepatide by reason of false designation of source and false and misleading advertising, the threat or effect of which is to destroy or substantially injure an industry in the United States, and based upon the importation into the United States, the sale for importation, and the sale within the United States after importation of certain products containing tirzepatide or purporting to contain tirzepatide by reason of infringement of U.S. Trademark No. 6,809,369. *Id.* The complaint also alleges that a domestic industry exists pursuant to subsection (a)(2) of section 337. *Id.*

The Commission's notice of investigation named as respondents Arctic Peptides LLC of Akeny, Iowa; Audrey Beauty Co. of Hong Kong, China; Biolabshop Limited of Lancaster, United Kingdom; Mew Mews Company Limited of Hong Kong, China; Strate Labs LLC of Spring, Texas; Steroide Kaufen of Bialystok, Poland; Super Human Store of Barcelona, Spain; Supopeptide of Cedar Grove, New Jersey; Triggered Supplements LLC of Clearwater, Florida; Unewlife of Cedar Grove, New Jersey; and Xiamen Austronext Trading Co., Ltd. of Fujian, China. *Id.* at 82915. The Office of Unfair Import Investigations (''OUII'') is also named as a party in this investigation. *Id.*

Respondents Unewlife, Supopeptide, and Steroide Kaufen were terminated pursuant to withdrawal of the complaint. *See* Order No. 8 (Mar. 7, 2024), *unreviewed by* Comm'n Notice (Mar. 21, 2024). Respondents Arctic Peptides LLC; Audrey Beauty Co., Ltd.; Biolabshop Limited; Mew Mews Co. Ltd.; Strate Labs LLC; Super Human Store; Triggered Supplements LLC (d/b/a The Triggered Brand); and Xiamen Austronext Trading Co., Ltd. (d/b/a AustroPeptide) have been found in default. *See* Order No. 13 (Apr. 22, 2024), *unreviewed by* Comm'n Notice (May 15, 2024).

On May 21, 2024, the complaint and notice of investigation were amended to add two respondents: Fibonacci Sequence LLC d/b/a GenX Peptides of Houston, Texas; and Paradigm Peptides of Michigan City, Indiana. Order No. 12 (Apr. 22, 2024), *unreviewed by* Comm'n Notice (May 21, 2024), 89 FR 46159–60 (May 28, 2024).

On May 8, 2024, Eli Lilly filed a motion to amend the complaint and notice of investigation to name an additional respondent, Total Compounding Pharmaceuticals of Australia. On that same date, Eli Lilly filed a motion for leave to serve Total Compounding Pharmaceuticals by alternative service via email, which was granted pursuant to Order No. 14 (May