

Irvine, California (collectively, “Complainants”). *See* 84 FR 8112–13 (Mar. 6, 2019). The complaint, as supplemented, alleges a violation of section 337 based upon the importation and sale in the United States of certain botulinum toxin products, processes for manufacturing or relating to same and certain products containing same by reason of misappropriation of trade secrets, the threat or effect of which is to destroy or substantially injure a domestic industry in the United States. *See id.* The notice of investigation names Daewoong and Evolus as respondents in this investigation. *See id.* The Office of Unfair Import Investigations is also a party to the investigation. *See id.*

On July 6, 2020, the Administrative Law Judge (“ALJ”) issued a final initial determination (“FID”) finding a violation of section 337 based on the importation and sale in the United States of Respondents’ botulinum neurotoxin products by reason of the misappropriation of trade secrets, the threat or effect of which is to destroy or substantially injure an industry in the United States. *See* FID at 273. The ALJ issued a recommended determination (“RD”) recommending that, if a violation is found, the Commission issue: (1) An LEO barring entry of certain botulinum toxin products that are imported and/or sold by respondents Daewoong and Evolus; and (2) a CDO against Evolus. The RD also recommends that the Commission impose a bond based on price differential during the period of Presidential review.

On July 28, 2020, the Commission issued a notice requesting statements on the public interest. *See* 85 FR 46711 (Aug. 3, 2020) (“the PI Notice”). On August 17–18, 2020, several non-parties filed submissions in response to the PI Notice.

On September 21, 2020, the Commission issued a notice determining to review the FID in part. *See* 85 FR 60489–90 (Sept. 25, 2020) (“the WTR/Remedy Notice”). Specifically, the Commission determined to review the FID’s findings with respect to subject matter jurisdiction, standing, trade secret existence and misappropriation, and domestic industry, including the existence of such domestic industry as well as any actual or threatened injury thereto. *See id.* The Commission determined not to review the remainder of the FID. *See id.* The Commission’s notice also requested written submissions on remedy, the public interest, and bonding. *See id.*

On October 9, 2020, the parties, including the IA, filed written submissions in response to the WTR/Remedy Notice, and on October 16, 2020, the parties filed responses to each other’s submissions. In addition, on October 5–9, 2020, several non-parties filed submissions on the proposed remedy and/or the public interest in response to the WTR/Remedy Notice.

Having examined the record of this investigation, including the FID, the RD, and the parties’ and non-parties’ submissions, the Commission has determined to affirm the FID in part and reverse in part. Specifically, as explained in the Commission Opinion filed concurrently herewith, the Commission has determined to affirm with modification the FID’s findings with respect to subject matter jurisdiction, standing, domestic industry as to BOTOX®, and trade secret existence and misappropriation as it relates to Medytox’s manufacturing processes. The Commission has also determined to reverse the FID’s finding that a trade secret exists with respect to Medytox’s bacterial strain. All findings in the FID that are not inconsistent with the Commission’s determination are affirmed.

Accordingly, the Commission finds that there is a violation of section 337. The Commission has determined that the appropriate remedy is an LEO against Respondents’ botulinum toxin products, and a CDO against Evolus, barring Respondents’ unfair acts for a duration of 21 months. The Commission has also determined that the public interest factors enumerated in subsections 337(d)(1) and (f)(1) (19 U.S.C. 1337(d)(1), (f)(1)) do not preclude the issuance of the LEO and CDO. The Commission has further determined to set a bond during the period of Presidential review in an amount of \$441 per 100U vial of Respondents’ accused products.

The Commission’s orders and opinion were delivered to the President and to the United States Trade Representative on the day of their issuance.

The investigation is terminated.

The Commission’s vote on this determination took place on December 16, 2020.

The authority for the Commission’s determination is contained in 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: December 16, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020–28158 Filed 12–21–20; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–649 and 731–TA–1523 (Final)]

Twist Ties From China; Scheduling of the Final Phase of Countervailing Duty and Antidumping Duty Investigations

AGENCY: United States International Trade Commission.

ACTION: Notice.

SUMMARY: The Commission hereby gives notice of the scheduling of the final phase of antidumping and countervailing duty investigation Nos. 701–TA–649 and 731–TA–1523 (Final) pursuant to the Tariff Act of 1930 (“the Act”) to determine whether an industry in the United States is materially injured or threatened with material injury, or the establishment of an industry in the United States is materially retarded, by reason of imports of twist ties from China, provided for in subheadings 8309.90.0000 and 5609.00.3000 of the Harmonized Tariff Schedule of the United States, preliminarily determined by the Department of Commerce (“Commerce”) to be subsidized and sold at less-than-fair-value.

DATES: December 3, 2020.

FOR FURTHER INFORMATION CONTACT: Christopher W. Robinson ((202) 205–2542), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission’s TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (<https://www.usitc.gov>). The public record for these investigations may be viewed on the Commission’s electronic docket (EDIS) at <https://edis.usitc.gov>.

SUPPLEMENTARY INFORMATION:

Scope.—For purposes of these investigations, Commerce has defined the subject merchandise as twist ties, which are thin, bendable ties for closing containers, such as bags, bundle items, or identifying objects. A twist tie in

most circumstances is comprised of one or more metal wires encased in a covering material, which allows the tie to retain its shape and bind against itself. However, it is possible to make a twist tie with plastic and no metal wires. The metal wire that is generally used in a twist tie is stainless or galvanized steel and typically measures between the gauges of 19 (.0410" diameter) and 31 (.0132") (American Standard Wire Gauge). A twist tie usually has a width between .075" and 1" in the cross-machine direction (width of the tie—measurement perpendicular with the wire); a thickness between .015" and .045" over the wire; and a thickness between .002" and .020" in areas without wire. The scope includes an all-plastic twist tie containing a plastic core as well as a plastic covering (the wing) over the core, just like paper and/or plastic in a metal tie. An all-plastic twist tie (without metal wire) would be of the same measurements as a twist tie containing one or more metal wires. Twist ties are commonly available individually in pre-cut lengths ("singles"), wound in large spools to be cut later by machine or hand, or in perforated sheets of spooled or single twist ties that are later slit by machine or by hand ("gangs").

The covering material of a twist tie may be paper (metallic or plain), or plastic, and can be dyed in a variety of colors with or without printing. A twist tie may have the same covering material on both sides or one side of paper and one side of plastic. When comprised of two sides of paper, the paper material is bound together with an adhesive or plastic. A twist tie may also have a tag or label attached to it or a pre-applied adhesive attached to it.

Excluded from the scope of the order are twist ties packaged with bags for sale together where the quantity of twist ties does not exceed twice the number of bags in each package. Also excluded are twist ties that constitute part of the packaging of the imported product, for example, merchandise anchored/secured to a backing with twist ties in the retail package or a bag of bread that is closed with a twist tie.

Twist ties are imported into the United States under Harmonized Tariff Schedule of the United States (HTSUS) subheadings 8309.90.00 and 5609.00.30. Subject merchandise may also be imported under HTSUS subheadings 3920.51.5000, 3923.90.0080, 3926.90.9990, 4811.59.6000, 4821.10.2000, 4821.10.4000, 4821.90.2000, 4821.90.4000, and 4823.90.8600. These HTSUS subheadings are provided for reference

only. The written description of the scope of the investigation is dispositive.

Background.—The final phase of these investigations is being scheduled pursuant to sections 705(b) and 731(b) of the Tariff Act of 1930 (19 U.S.C. 1671d(b) and 1673d(b)), as a result of affirmative preliminary determinations by Commerce that certain benefits which constitute subsidies within the meaning of § 703 of the Act (19 U.S.C. 1671b) are being provided to manufacturers, producers, or exporters in China of twist ties, and that such products are being sold in the United States at less than fair value within the meaning of § 733 of the Act (19 U.S.C. 1673b). The investigations were requested in petitions filed on June 26, 2020, by Bedford Industries Inc., Worthington, Minnesota.

For further information concerning the conduct of this phase of the investigations, hearing procedures, and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A and B (19 CFR part 201), and part 207, subparts A and C (19 CFR part 207).

Participation in the investigations and public service list.—Persons, including industrial users of the subject merchandise and, if the merchandise is sold at the retail level, representative consumer organizations, wishing to participate in the final phase of these investigations as parties must file an entry of appearance with the Secretary to the Commission, as provided in § 201.11 of the Commission's rules, no later than 21 days prior to the hearing date specified in this notice. A party that filed a notice of appearance during the preliminary phase of the investigations need not file an additional notice of appearance during this final phase. The Secretary will maintain a public service list containing the names and addresses of all persons, or their representatives, who are parties to the investigations.

Please note the Secretary's Office will accept only electronic filings during this time. Filings must be made through the Commission's Electronic Document Information System (EDIS, <https://edis.usitc.gov>). No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice.

Limited disclosure of business proprietary information (BPI) under an administrative protective order (APO) and BPI service list.—Pursuant to § 207.7(a) of the Commission's rules, the Secretary will make BPI gathered in the final phase of these investigations available to authorized applicants under the APO issued in the investigations,

provided that the application is made no later than 21 days prior to the hearing date specified in this notice. Authorized applicants must represent interested parties, as defined by 19 U.S.C. 1677(9), who are parties to the investigations. A party granted access to BPI in the preliminary phase of the investigations need not reapply for such access. A separate service list will be maintained by the Secretary for those parties authorized to receive BPI under the APO.

Staff report.—The prehearing staff report in the final phase of these investigations will be placed in the nonpublic record on February 2, 2021, and a public version will be issued thereafter, pursuant to § 207.22 of the Commission's rules.

Hearing.—The Commission will hold a hearing in connection with the final phase of these investigations beginning at 9:30 a.m. on February 16, 2021. Information about the place and form of the hearing, including about how to participate in and/or view the hearing, will be posted on the Commission's website at <https://www.usitc.gov/calendarpad/calendar.html>. Interested parties should check the Commission's website periodically for updates. Requests to appear at the hearing should be filed in writing with the Secretary to the Commission on or before February 9, 2021. A nonparty who has testimony that may aid the Commission's deliberations may request permission to present a short statement at the hearing. All parties and nonparties desiring to appear at the hearing and make oral presentations should attend a prehearing conference to be held at 9:30 a.m. on February 11, 2021. Oral testimony and written materials to be submitted at the public hearing are governed by sections 201.6(b)(2), 201.13(f), and 207.24 of the Commission's rules. Parties must submit any request to present a portion of their hearing testimony *in camera* no later than 7 business days prior to the date of the hearing.

Written submissions.—Each party who is an interested party shall submit a prehearing brief to the Commission. Prehearing briefs must conform with the provisions of § 207.23 of the Commission's rules; the deadline for filing is February 9, 2021. Parties may also file written testimony in connection with their presentation at the hearing, as provided in § 207.24 of the Commission's rules, and posthearing briefs, which must conform with the provisions of § 207.25 of the Commission's rules. The deadline for filing posthearing briefs is February 25, 2021. In addition, any person who has

not entered an appearance as a party to the investigations may submit a written statement of information pertinent to the subject of the investigations, including statements of support or opposition to the petition, on or before February 25, 2021. On March 16, 2021, the Commission will make available to parties all information on which they have not had an opportunity to comment. Parties may submit final comments on this information on or before March 18, 2021, but such final comments must not contain new factual information and must otherwise comply with § 207.30 of the Commission's rules. All written submissions must conform with the provisions of § 201.8 of the Commission's rules; any submissions that contain BPI must also conform with the requirements of §§ 201.6, 207.3, and 207.7 of the Commission's rules. The Commission's *Handbook on Filing Procedures*, available on the Commission's website at https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf, elaborates upon the Commission's procedures with respect to filings.

Additional written submissions to the Commission, including requests pursuant to § 201.12 of the Commission's rules, shall not be accepted unless good cause is shown for accepting such submissions, or unless the submission is pursuant to a specific request by a Commissioner or Commission staff.

In accordance with §§ 201.16(c) and 207.3 of the Commission's rules, each document filed by a party to the investigations must be served on all other parties to the investigations (as identified by either the public or BPI service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

Authority: These investigations are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.21 of the Commission's rules.

By order of the Commission.

Issued: December 16, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020–28140 Filed 12–21–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—CHEDE–8

Notice is hereby given that, on December 15, 2020, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), CHEDE–8 (“CHEDE–8”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Caterpillar, Inc., Peoria, IL, has been added as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and CHEDE–8 intends to file additional written notifications disclosing all changes in membership.

On December 4, 2019, CHEDE–8 filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on December 30, 2019 (84 FR 71977).

The last notification was filed with the Department on October 20, 2020. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on November 19, 2020 (85 FR 73751).

Suzanne Morris,

Chief, Premerger and Division Statistics, Antitrust Division.

[FR Doc. 2020–28141 Filed 12–21–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—IMS Global Learning Consortium, Inc.

Notice is hereby given that, on December 3, 2020, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), IMS Global Learning Consortium, Inc. (“IMS Global”) has filed written notifications simultaneously with the Attorney

General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Australian Council for Educational Research, Camberwell, AUSTRALIA; ClassEDU Inc., Raleigh, NC; GreenLight Credentials, Dallas, TX; Magic Software Inc., New York, NY; Manabie International Pte Ltd, Singapore, SINGAPORE; Richland School District No. 2, Columbia, SC; and Virtual Virginia, Floyd, VA, have been added as parties to this venture.

Also, IBM, Cambridge, MA; Paradigm, Inc., Virginia Beach, VA; Badgewell, Giza, EGYPT; VidGrid, St. Paul, MN; and Unox Portal (Pragnya Technologies Pty), Sydney, AUSTRALIA, have withdrawn as parties to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and IMS Global intends to file additional written notifications disclosing all changes in membership.

On April 7, 2000, IMS Global filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on September 13, 2000 (65 FR 55283).

The last notification was filed with the Department on September 16, 2020. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on October 15, 2020 (85 FR 65426).

Suzanne Morris,

Chief, Premerger and Division Statistics, Antitrust Division.

[FR Doc. 2020–28138 Filed 12–21–20; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–758]

Importer of Controlled Substances Application: Fresenius Kabi USA, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Fresenius Kabi USA, LLC has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTAL INFORMATION** listed below for further drug information.