submitted by the named respondents in accordance with section 210.13 of the Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: February 28, 2019.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2019–03988 Filed 3–5–19; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1145]

Certain Botulinum Toxin Products, Processes for Manufacturing or Relating to Same and Certain Products Containing Same; Institution of Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that a complaint was filed with the U.S. International Trade Commission on January 30, 2019, under section 337 of the Tariff Act of 1930, as amended, on behalf of Medytox Inc. of South Korea; Allergan plc of Ireland; Allergan, Inc., Irvine, California. Supplements to the complaint were filed on February 12, 2019, February 13, 2019, and February 14, 2019. The complaint alleges violations of section 337 based upon the importation into the United States, the sale for importation, and the sale within the United States after importation 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.

The complainants request that the Commission institute an investigation and, after the investigation, issue a limited exclusion order and cease and desist orders.

ADDRESSES: The complaint, except for any confidential information contained therein, is available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW, Room 112, Washington, DC 20436, telephone (202) 205–2000. Hearing impaired individuals are advised that information on this matter can be obtained 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 at *https://www.usitc.gov*. The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

FOR FURTHER INFORMATION CONTACT: Pathenia M. Proctor, The Office of Unfair Import Investigations, U.S. International Trade Commission, telephone (202) 205–2560.

SUPPLEMENTARY INFORMATION:

Authority: The authority for institution of this investigation is contained in section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, and in section 210.10 of the Commission's Rules of Practice and Procedure, 19 CFR 210.10 (2018).

Scope of Investigation: Having considered the complaint, the U.S. International Trade Commission, on February 28, 2019, *ordered that*—

(1) Pursuant to section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine whether there is a violation of subsection (a)(1)(A) of section 337 in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain products identified in paragraph (2) 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;

(2) Pursuant to section 210.10(b)(1) of the Commission's Rules of Practice and Procedure, 19 CFR 210.10(b)(1), the plain language description of the accused products or category of accused products, which defines the scope of the investigation, is botulinum neurotoxin products manufactured by Daewoong Pharmaceuticals Co., Ltd., specifically: (1) DWP–450 (prabotulinumtoxinA), variously marketed under the brand names Nabota®, Jeuveau™ and other brand names; (2) products containing or derived from DWP–450; and (3) products containing or derived from the BTX strain assigned the high-risk pathogen control number 4–029–CBB–IS–001 by the Korean Centers for Disease Control and Prevention or the manufacturing process used to manufacture DWP–450;

(3) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

(a) The complainants are: Medytox Inc., 626 Tehran Road, Gangnam, Seoul, South Korea; Allergan plc, Clonshaugh Business and Technology Park, Coolock, Dublin, D17 E400, Irland; Allergan, Inc., 2525 Dupont Drive, Irvine, CA 92612.

(b) The respondents are the following entities alleged to be in violation of section 337, and are the parties upon which the complaint is to be served: Daewoong Pharmaceuticals Co., Ltd., Bongeunsaro 114-gil 12, Gangnam, Seoul, 06170; South Korea; Evolus, Inc., 17901 Von Karman Avenue, Suite 150, Irvine, CA 92614.

(c) The Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street SW, Suite 401, Washington, DC 20436; and

(4) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with section 210.13 of the Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be

deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: February 28, 2019.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2019–03987 Filed 3–5–19; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—X12 Incorporated

Notice is hereby given that, on February 11, 2019, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), X12 Incorporated (''X12'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the name and principal place of business of the standards development organization and (2) the nature and scope of its standards development activities. The notifications were filed for the purpose of invoking the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the name and principal place of business of the standards development organization is: X12 Incorporated, McClean, VA. The nature and scope of X12's standards development activities are: The development and maintenance of cross industry e-commerce standards that improve business process interoperability and facilitate business information exchange supporting the finance, government, supply chain, transportation and insurance industries and associated business partners.

Suzanne Morris,

Chief, Premerger and Division Statistics Unit, Antitrust Division.

[FR Doc. 2019–04010 Filed 3–5–19; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—HEI Industry Group

Notice is hereby given that, on January 28, 2019, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), HEI Industry Group (''HIG'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the identities of the parties to the venture and (2) the nature and objectives of the venture. The notifications were filed for the purpose of invoking the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the identity of the parties to the venture are: Chevron North America Exploration and Production Company, a division of Chevron U.S.A. Inc., Houston, TX; ConocoPhillips Company, Houston, TX; Exxon Mobil Corporation, Irving, TX; Halliburton Energy Services, Inc., Houston, TX; Noble Energy, Inc., Houston TX, SWEPI LP (SWEPI), Houston, TX; Schlumberger Technology Corporation, Houston, TX; BHP, Houston, TX; Statoil Gulf Services LLC, Houston, TX; and Schlumberger Limited, N.V., Houston, TX. The general area of HIG's planned activity is to commence a joint industry-government research initiative entitled HEI's Energy Research Program to (1) evaluate the existing health and exposure literature related to potential impacts from onshore oil and natural gas operations; and possibly (2) conduct a study to assess potential exposures from those operations. The industry sponsors have created the HEI Industry Group (HIG) to facilitate coordinated input to HEI. HEI is a nonprofit organization chartered in 1980 as an independent research institute to provide high-quality, impartial, and relevant science on the health effects of air pollution. The HEI-managed program represents a first-of-its-kind, comprehensive collaboration between the oil and gas industry and government to assess exposure to chemical stressors associated with onshore unconventional oil and natural gas operations. Part 1 of the research program will last for approximately one year and will evaluate the existing health and exposure literature as well as conduct workshops to inform the literature reviews and frame research needs. Part 2 (an exposure study) is being considered and will be informed by the results of the Part 1 literature review. If Part 2 goes forward, future studies will be considered as warranted by the Part 2 results.

Suzanne Morris,

Chief, Premerger and Division Statistics Unit, Antitrust Division.

[FR Doc. 2019–04009 Filed 3–5–19; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: The registrant listed below has applied for and been granted registration by-the Drug Enforcement Administration (DEA) as an importer of schedule I controlled substances.

SUPPLEMENTARY INFORMATION: The company listed below applied to be registered as an importer of scheduled I controlled substances. Information on the previously published notice is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for this notice.

Company	FR Docket	Published
Agilent Technologies	83 FR 66751	December 27, 2018.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrant to import the applicable basic classes of schedule I controlled substances is consistent with the public interest and with United States obligations under