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 amended complaint, the U.S. International Trade Commission, on December 6, 2018, *ordered that*—

(1) Pursuant to subsection (b) of 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)(B) 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 infringement of one or more of claims 1, 2, 4, 5, 7, 8, 10, 12, 13, 16, 17, 20, and 21 of the '669 patent; claims 1–4, 9–11, 13, 14, 19–21, 24, 28, and 29 of the '139 patent; claims 1–3, 5–9, 12, and 17–20 of the '568 patent; claims 1, 2, 4–6, 8–10, 16, 19, 21, and 27 of the '130 patent; and claims 1–4, 6, 9, 11, 12, 18–23, and 27 of the '915 patent; and whether an industry in the United States exists as required by subsection (a)(2) of section 337;

(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 ''nicotine vaporizer devices and the associated pods sold for use with the devices, and components thereof'';

(3) Pursuant to Commission Rule 210.50(b)(1), 19 CFR 210.50(b)(1), the presiding administrative law judge shall take evidence or other information and hear arguments from the parties or other interested persons with respect to the public interest in this investigation, as appropriate, and provide the Commission with findings of fact and a recommended determination on this issue, which shall be limited to the statutory public interest factors set forth in 19 U.S.C. 1337(d)(1), (f)(1), (g)(1);

(4) 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: Juul Labs, Inc., 560 20th Street, San Francisco, CA 94107.

(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:

J Well France S.A.S., 50 rue de Miromesnil, 75008 Paris, France
Bo Vaping, 591 Stewart Avenue, Garden City, NY 11530
MMS Distribution LLC, 195 Lake Louise Marie Road, Rock Hill, NY 12775
The Electric Tobacconist, LLC, 3235 Prairie Avenue, Boulder, CO 80301
Vapor 4 Life Holdings, Inc., 4080 Commercial Avenue, Suite A, Northbrook, IL 60062
Eonsmoke, LLC, 1500 Main Ave, 2nd Floor, Clifton, NJ 07011
ZLab S.A., Ave. Golero, 911 Office 27, Punta del Este—Maldonado—Uruguay 20100
Ziip Lab Co., Limited, E district 4F, 5 building, Wen Ge Industrial Zone, Heshuikou Gongming St., Guangming New District, Shenzhen City, Guangdong Province, China 518106
Shenzhen Yibo Technology Co., Ltd., E district 4F, 5 building, Wen Ge Industrial Zone, Heshuikou, Gongming St., Guangming New District, Shenzhen City, Guangdong Province, China 518106
XFire, Inc., 820 Summer Park Dr., Suite 700, Stafford, TX 77477
ALD Group Limited, No. 2, 3rd Industrial Road, Shixin Community, Shiyan Street, Bao'an District, Shenzhen City, Guangdong Province, China 518108
Flair Vapor LLC, 2500 Hamilton Blvd., Suite B, South Plainfield, NJ 07080
Shenzhen Joecig Technology Co., Ltd., 1F–5F, Building 17, Quarter G ShaJing Rd., Gonghe 3rd Industry District, Baoan District, Shenzhen City, Guangdong Province, China 518104
Myle Vape Inc., 8085 Chevy Chase Street, Jamaica, NY 11432
Vapor Hub International, Inc., 1871 Tapo Street, Simi Valley, CA 93063
Limitless Mod Co., 4590 Ish Drive, Suite 100, Simi Valley, CA 93063
Asher Dynamics, Inc., 14345 Pipeline Avenue, Chino, CA 91710
Ply Rock, 14345 Pipeline Avenue, Chino, CA 91710
Infinite-N Technology Limited, 4F, iTone Digital Park, Xin Fa San Road, Sha Jing Shenzhen City, Guangdong Province, China 518200
King Distribution LLC, 281 Route 46 West, Elmwood Park, NJ 07407
Keep Vapor Electronic Tech. Co., Ltd., Block D, XinLong Techno Park, ShaJing Town, Bao An District, Shenzhen, China

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

(5) 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: December 10, 2018.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2018–26995 Filed 12–12–18; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

Notice of Appointment of Individuals To Serve as Members of the Performance Review Board

AGENCY: United States International Trade Commission.

ACTION: Notice.

DATES: December 7, 2018.

FOR FURTHER INFORMATION CONTACT: Eric Mozie, Director of Human Resources, or Ronald Johnson, Senior Human Resources Specialist, U.S. International Trade Commission (202) 205–2651.

SUPPLEMENTARY INFORMATION: The Chairman of the U.S. International Trade Commission has appointed the following individuals to serve on the Commission's Performance Review Board (PRB):

Chair of the PRB: Commissioner Irving A. Williamson
Vice-Chair of the PRB: Commissioner Meredith Broadbent
Member—John Ascienzo
Member—Dominic Bianchi
Member—Nannette Christ
Member—Catherine DeFilippo
Member—James Holbein
Member—Margaret Macdonald
Member—Stephen A. McLaughlin
Member—William Powers

Authority: This notice is published in the **Federal Register** pursuant to the requirement of 5 U.S.C. 4314(c)(4). Hearing impaired individuals are advised that information on this matter can be obtained by contacting our TDD terminal on (202) 205–1810.

By order of the Commission.

Issued: December 4, 2018.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2018–26742 Filed 12–12–18; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Siegfried USA, LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 14, 2019. Such persons may also file a written request for a hearing on the application on or before January 14, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (''Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on November 05, 2018, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Opium, raw	9600	II
Poppy Straw Concentrate	9670	II

The company plans to import the listed controlled substances to manufacture bulk active pharmaceutical ingredients (API) for distribution to its customers.

Dated: December 3, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–27032 Filed 12–12–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Mylan Pharmaceuticals Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 14, 2019. Such persons may also file a written request for a hearing on the application on or before January 14, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (''Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on October 10, 2018, Mylan Pharmaceuticals Inc., 2898 Manufacturers Road, Greensboro, North Carolina 27406–4600 applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Nabilone	7379	II

The company plans to import the FDA approved drug product in finished dosage form for distribution to its customers. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2).