in the National Register of Historic Places. Nominations for their consideration were received by the National Park Service before October 10, 2020. Pursuant to Section 60.13 of 36 CFR part 60, comments are being accepted concerning the significance of the nominated properties under the National Register criteria for evaluation.

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.

Nominations submitted by State or Tribal Historic Preservation Officers:

ALABAMA

Mobile County

Midtown Historic District (Boundary Increase), 2401–2403 and 2407 Old Shell Rd., Mobile, BC100005805

CALIFORNIA

Los Angeles County

Pasadena Field Archery Range, 415 South Arroyo Blvd., Pasadena, SG100005799

Riverside County

Desert Golf Course, 301 North Belardo Rd., Palm Springs, SG100005813

San Francisco County

Whelan, John A., House, 1315 Waller St., San Francisco, SG100005794

COLORADO

Denver County

First Avenue Hotel, 101 North Broadway, Denver, SG100005800

CONNECTICUT

Litchfield County

Winsted Water Works,Winchester Rd. (north side), Old Waterbury Tpk./Rugg Brook Rd., Winchester, SG100005797

GEORGIA

Fulton County

Cascade Heights Commercial Historic District, Centered on the jct. of Cascade Rd. SW and Benjamin E. Hayes Dr. SW, Atlanta, SG100005817

IOWA

Wapello County

Agassiz School, 608 East Williams St., Ottumwa, SG100005787

MASSACHUSETTS

Suffolk County

Crawford Street Historic District, 5–38 Crawford St., 42 Elm Hill Ave., 621 Warren St., Boston, SG100005798

NEW JERSEY

Mercer County

V. Henry Rothschild-F.A. Straus and Co. Atlantic Products Corporation Mill Complex, 1 North Johnston Ave., Hamilton Township, SG100005815

OHIO

Stark County

St. Joseph Roman Catholic Church Complex, 2427 Tuscarawas St. West, Canton, SG100005806

PUERTO RICO

San Juan Municipality

Rafael Cordero Graded School, (Early Twentieth Century Schools in Puerto Rico TR), Calle Aurora Esq. Horae Prada 15, Santurce vicinity, MP100005816

SOUTH DAKOTA

Roberts County

Sisseton School, (Schools in South Dakota MPS), 302 East Maple St., Sisseton, MP100005818

Sisseton School, (Federal Relief Construction in South Dakota MPS), 302 East Maple St., Sisseton, MP100005818

TENNESSEE

Blount County

Millennium Manor, 500 North Wright Rd., Alcoa, SG100005788

Montgomery County

Mt. Olive Cemetery, 951 Cumberland Dr., Clarksville, SG100005789

Rhea County

First Avenue Methodist Episcopal Church, 240 1st Ave., Dayton, SG100005790

Sullivan County

Kingsport Hosiery Mills, 435 Press St., Kingsport, SG100005791

Washington County

Johnson City Postal Savings Bank and Post Office, 401 Ashe St., Johnson City, SG100005792

Wayne County

Hughes House, 204 West Pillow St., Clifton, SG100005793

VIRGINIA

Alexandria Independent City

George Washington High School, 1005 Mount Vernon Ave., Alexandria, SG100005803

Craig County

Bellevue, 14505 Cumberland Gap Rd. (VA 42), New Castle vicinity, SG100005801

Halifax County

Oak Cliff, 10000 Huell Matthews Hwy. (US 501), Alton vicinity, SG100005804

Washington County

Depot Square Historic District, Wall St. South, Depot Sq. SW, Front St. SW, Grand St. SW, Abingdon, SG100005802

Additional documentation has been received for the following resources:

ARIZONA

Pima County

Armory Park Historic Residential District (Additional Documentation), East 12th St. to 19th St., Stone Ave. to 2nd Ave., Tucson, AD76000378

Hughes, Sam, Neighborhood Historic District (Additional Documentation), Roughly bounded by East Speedway Blvd., North Campbell Ave., East 7th St. and North, Bentley Ave., Tucson, AD94001164

KANSAS

Trego County

Wilcox School-District 29 & District 14 (Additional Documentation), (Public Schools of Kansas MPS), Rural Route –15 mi. south of WaKeeney on KS 283, Ransom, AD06000393

(Authority: Section 60.13 of 36 CFR part 60)

Dated: October 14, 2020.

Sherry A. Frear,

Chief, National Register of Historic Places/National Historic Landmarks Program.

[FR Doc. 2020–23681 Filed 10–26–20; 8:45 am]

BILLING CODE 4312–52–P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 731–TA–1229–1230 (Review)]

Monosodium Glutamate From China and Indonesia

Determination

On the basis of the record [1] developed in the subject five-year reviews, the United States International Trade Commission (''Commission'') determines, pursuant to the Tariff Act of 1930 (''the Act''), that revocation of the antidumping duty orders on monosodium glutamate (''MSG'') from China and Indonesia would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.

Background

The Commission instituted these reviews on October 1, 2019 (84 FR 52129) and determined on January 6, 2020 that it would conduct full reviews (85 FR 3421, January 21, 2020). Notice of the scheduling of the Commission's reviews and of a public hearing to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the

[1] The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

notice in the **Federal Register** on May 13, 2020 (85 FR 28663). In light of the restrictions on access to the Commission building due to the COVID–19 pandemic, the Commission conducted its hearing through written testimony and video conference on August 25, 2020. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). The Commission determined that these reviews were extraordinarily complicated and extended the review period by up to 90 days. It completed and filed its determinations in these reviews on October 21, 2020. The views of the Commission are contained in USITC Publication 5127 (October 2020), entitled *Monosodium Glutamate from China and Indonesia: Investigation Nos. 731–TA–1229–1230 (Review).*

By order of the Commission.

Issued: October 21, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020–23696 Filed 10–26–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–736]

Importer of Controlled Substances Application: Mylan Pharmaceuticals

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Mylan Pharmaceuticals 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.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before November 27, 2020. Such persons may also file a written request for a hearing on the application on or before November 27, 2020.

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 a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a 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: In accordance with 21 CFR 1301.34(a), this is notice that on October 7, 2020, Mylan Pharmaceuticals, Incorporated, 3711 Collins Ferry Road, Morgantown, West Virginia 26505–2362, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Amphetamine	1100	II
Methylphenidate	1724	II
Oxycodone	9143	II
Hydromorphone	9150	II
Methadone	9250	II
Morphine	9300	II
Fentanyl	9801	II

The company plans to import finished dosage forms for analytical testing and distribution for clinical trials. No other activity for these drug codes is authorized for this registration.

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). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–23766 Filed 10–26–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–738]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: API GLOBAL LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: The Drug Enforcement Administration (DEA) is providing notice of an application it has received from an entity applying to be registered to manufacture in bulk basic class(es) of controlled substances listed in schedule I. DEA intends to evaluate this and other pending applications according to proposed regulations that, if finalized, would govern the program of growing marihuana for scientific and medical research under DEA registration.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections to the issuance of the proposed registration on or before December 28, 2020.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW 8701 Morrissette Drive, Springfield, Virginia 22152. To ensure proper handling of comments, please reference Docket No -. DEA–738 in all correspondence, including attachments.

SUPPLEMENTARY INFORMATION: The Controlled Substances Act (CSA) prohibits the cultivation and distribution of marihuana except by persons who are registered under the CSA to do so for lawful purposes. In accordance with the purposes specified in 21 CFR 1301.33(a), DEA is providing notice that the entity identified below has applied for registration as a bulk manufacturer of schedule I controlled substances. In response, registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections of the requested registration, as provided in this notice. This notice does not constitute any evaluation or determination of the merits of the application submitted.

The applicant plans to manufacture bulk active pharmaceutical ingredients (APIs) for product development and distribution to DEA registered researchers. If the application for registration is granted, the registrant would not be authorized to conduct other activity under this registration aside from those coincident activities specifically authorized by DEA regulations. DEA will evaluate the application for registration as a bulk manufacturer for compliance with all applicable laws, treaties, and regulations and to ensure adequate safeguards against diversion are in place.

As this applicant has applied to become registered as a bulk manufacturer of marihuana, the application will be evaluated under the criteria of 21 U.S.C. 823(a). DEA proposes to conduct this evaluation in the manner described in the rule proposed at 85 FR 16292, published on March 23, 2020, if finalized.