

at the retail level, representative consumer organizations have the right to appear as parties in Commission antidumping and countervailing duty investigations. The Secretary will prepare a public service list containing the names and addresses of all persons, or their representatives, who are parties to the investigations. As provided in section 207.20 of the Commission's rules, the Director of the Office of Investigations will circulate draft questionnaires for the final phase of the investigations to parties to the investigations, placing copies on the Commission's Electronic Document Information System (EDIS, <https://edis.usitc.gov>), for comment.

### Background

On February 26, 2025, the U.S. Chassis Manufacturers Coalition, whose members are Cheetah Chassis Corporation, Berwick, Pennsylvania and Stoughton Trailers LLC, Stoughton, Wisconsin, filed petitions with the Commission and Commerce, alleging that an industry in the United States is materially injured or threatened with material injury by reason of subsidized imports of chassis and subassemblies from Mexico and Thailand and LTFV imports of chassis and subassemblies from Mexico, Thailand, and Vietnam. Accordingly, effective February 26, 2025, the Commission instituted countervailing duty investigation Nos. 701-TA-755-756 and antidumping duty investigation Nos. 731-TA-1734-1736 (Preliminary).

Notice of the institution of the Commission's investigations and of a public conference 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 notice in the **Federal Register** of March 4, 2025 (90 FR 11180). The Commission conducted its conference on March 19, 2025. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to §§ 703(a) and 733(a) of the Act (19 U.S.C. 1671b(a) and 1673b(a)). It completed and filed its determinations in these investigations on April 14, 2025. The views of the Commission are contained in USITC Publication 5612 (April 2025), entitled *Chassis and Subassemblies from Mexico, Thailand, and Vietnam: Investigation Nos. 701-TA-755-756 and 731-TA-1734-1736 (Preliminary)*.

By order of the Commission.

Issued: April 14, 2025.

**Lisa Barton,**

*Secretary to the Commission.*

[FR Doc. 2025-06672 Filed 4-17-25; 8:45 am]

**BILLING CODE 7020-02-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1527]

#### Importer of Controlled Substances Application: Scottsdale Research Institute

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Scottsdale Research Institute has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before May 19, 2025. Such persons may also file a written request for a hearing on the application on or before May 19, 2025.

**ADDRESSES:** The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitted comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152, and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on January 13, 2025, Scottsdale Research Institute, 12815 North Cave Creek Road, Phoenix, Arizona 85022, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana .....	7360	I

The company plans to import Active Pharmaceutical Ingredients for clinical research by Schedule I registrants. The company aim to meet the needs of Schedule I researchers with these novel compounds. In reference to drug codes 7360 (Marihuana), the company plans to import a synthetic cannabidiol. No other activity for this drug code 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.

**Matthew Strait,**

*Deputy Assistant Administrator.*

[FR Doc. 2025-06705 Filed 4-17-25; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1529]

#### Bulk Manufacturer of Controlled Substances Application: Patheon API Services Inc.

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Patheon API Services Inc. has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before June 17, 2025. Such persons may also file a written request for a hearing on the application on or before June 17, 2025.

**ADDRESSES:** The Drug Enforcement Administration requires that all

comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on February 5, 2025, Patheon API Services Inc., 101 Technology Place, Florence, South Carolina 29501, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Dimethyltryptamine	7435	I
Psilocybin .....	7437	I
Psilocyn .....	7438	I
Amphetamine .....	1100	II
Methadone .....	9250	II

The company plans to bulk manufacture the listed controlled substances to support the clinical trials of their customers. No other activities for these drug codes are authorized for this registration.

**Matthew Strait,**  
Deputy Assistant Administrator.

[FR Doc. 2025-06707 Filed 4-17-25; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1532]

#### Importer of Controlled Substances Application: Myonex LLC

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Myonex LLC has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and

applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before May 19, 2025. Such persons may also file a written request for a hearing on the application on or before May 19, 2025.

**ADDRESSES:** The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on January 9, 2025, Myonex LLC, 100 Progress Drive, Horsham, Pennsylvania 19044, 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
Lisdexamfetamine ..	1205	II
Methylphenidate .....	1724	II
Nabilone .....	7379	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Hydrocodone .....	9193	II
Morphine .....	9300	II
Oxymorphone .....	9652	II
Fentanyl .....	9801	II

The company plans to import the listed controlled substances in dosage form for clinical trials, research, and analytical purposes. No other activities for these drug codes are 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.

**Matthew Strait,**

Deputy Assistant Administrator.

[FR Doc. 2025-06712 Filed 4-17-25; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1513]

#### Bulk Manufacturer of Controlled Substances Application: Stepan Company

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Stepan Company has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before June 17, 2025. Such persons may also file a written request for a hearing on the application on or before June 17, 2025.

**ADDRESSES:** The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on January 31, 2025, Stepan Company, 100 West Hunter Avenue, Maywood, New Jersey 07607-