

is notice that on February 9, 2026, Pisgah Laboratories Inc, 3222 Old Hendersonville Highway, Pisgah Forest, North Carolina 28768, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
4-Bromo-2,5-dimethoxy-phenethylamine	7392	I
Psilocybin	7437	I
Methylone (3,4-Methylenedioxy-N-methylcathinone)	7540	I
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Cocaine	9041	II
Diphenoxylate	9170	II
Meperidine	9230	II
Methadone	9250	II
Tapentadol	9780	II

The company plans to bulk manufacture the above-listed controlled substances in bulk for internal research purposes and distribution to its customers. No other activities for these drug codes are authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2026-04971 Filed 3-12-26; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1676]

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 May 12, 2026. Such persons may also file a written request for a hearing on the application on or before May 12, 2026.

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 2, 2026, Stepan Company, 100 West Hunter Avenue, Maywood, New Jersey 07607-1021, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Cocaine	9041	II
Ecgonine	9180	II

The company plans to bulk manufacture the listed controlled substances for use as internal intermediates or for sale to its customers. No other activities for these drug codes are authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2026-04969 Filed 3-12-26; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1679]

Importer of Controlled Substances Application: Lonza Tampa, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Lonza Tampa, 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 April 13, 2026. Such persons may also file a written request

for a hearing on the application on or before April 13, 2026.

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 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 February 17, 2026, Lonza Tampa, LLC, 4901 West Grace Street, Tampa, Florida 33607-3805, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Psilocybin	7437	I

The company plans to import drug code 7437 (Psilocybin) as bulk active pharmaceutical ingredient and as finished dosage units for clinical trials, research, and analytical purposes. 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.

Thomas Prevoznik,
Deputy Assistant Administrator.
[FR Doc. 2026-04972 Filed 3-12-26; 8:45 am]
BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1681]

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 May 12, 2026. Such persons may also file a written request for a hearing on the application on or before May 12, 2026.

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 19, 2026, 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 for distribution to its customers. No other activities for these drug codes are authorized for this registration.

Thomas Prevoznik,
Deputy Assistant Administrator.
[FR Doc. 2026-04973 Filed 3-12-26; 8:45 am]
BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Consent Decree Under the Comprehensive Environmental Response Compensation and Liability Act

On March 6, 2026, the Department of Justice lodged a proposed Consent Decree with the United States District Court for the Middle District of Pennsylvania in the lawsuit entitled *United States of America and the Commonwealth of Pennsylvania v. Rutgers Organics LLC*, Civil Action No. 4:26-cv-00560-MWB.

The lawsuit was initiated by a complaint filed by the United States and the Commonwealth of Pennsylvania (the "Trustees") in their capacity as the legally designated trustees for natural resources in Pennsylvania. The complaint alleged, inter alia, that the Defendant was liable for damages for injury to, destruction of, or loss of natural resources resulting from the release of hazardous substances at and from the Centre County Kepone Site (the "Site") in College Township, Centre County, Pennsylvania, pursuant to the federal Comprehensive Environmental Response, Compensation and Liability Act of 1980 and the Pennsylvania Hazardous Site Cleanup Act.

The Consent Decree resolves the claims of the Trustees against the Defendant for a total payment of \$795,263. Of this amount, \$618,060 will be paid into the United States' Natural Resource Damages Assessment and Restoration Fund managed by the U.S. Department of the Interior, which will reimburse the Department's natural resource damages assessment activities and fund projects aimed at restoring the injured natural resources at the Site. The remaining \$177,203 will be paid to

the Commonwealth of Pennsylvania to reimburse the Commonwealth's natural resource damages assessment costs. In addition, under Commonwealth oversight the Defendant will construct a parking lot and a trail to facilitate public fishing access in Spring Creek at a location near the Site, and it will ensure the recordation of a conservation easement on property adjoining Spring Creek to facilitate water quality restoration efforts. In return for these payments and restoration activities, the United States and the Commonwealth will confer on the Defendant covenants not to sue for natural resource damages known or reasonably ascertainable as of the date of lodging of the Consent Decree.

In accordance with CERCLA, the Trustees have also written a draft Restoration Plan that described proposed alternatives for restoring the natural resources and natural resource services injured by the release of hazardous substances from the Site. The restoration alternatives evaluated in the Restoration Plan are (1) No Action/Natural Recovery; (2) Conservation Easement and Angler Access at NRDAR Property; (3) Future Repair and Maintenance of Existing In-Stream Structures along NRDAR Property; (4) Additional Land Conservation In Spring Creek Watershed; and (5) Additional Restoration Projects within Spring Creek Watershed.

The publication of this notice opens a period for public comment on the proposed Consent Decree and the draft Restoration Plan.

Comments on the proposed Consent Decree should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States of America and the Commonwealth of Pennsylvania v. Rutgers Organics LLC*, D.J. Ref. No. 90-11-3-1436/4. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

To submit comments:	Send them to:
By email	pubcomment-ees.enrd@usdoj.gov .
By mail	Assistant Attorney General, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

During the public comment period, the proposed Consent Decree may be examined and downloaded at this Justice Department website: <https://www.justice.gov/enrd/consent-decrees>.