

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]

**BILLING CODE P**

**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-