

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1684]

**Importer of Controlled Substances Application: United States Pharmacopeial Convention**

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

**ACTION:** Notice of application.

**SUMMARY:** United States Pharmacopeial Convention 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 20, 2026. Such persons may also file a written request for a hearing on the application on or before April 20, 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 11, 2026, United States Pharmacopeial Convention, 7135 English Muffin Way, Frederick, Maryland 21704, applied to be registered as an importer of the

following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Cathinone .....	1235	I
Methcathinone .....	1237	I
Methaqualone .....	2565	I
Lysergic acid diethylamide.	7315	I
4-Methyl-2,5-dimethoxyamphetamine.	7395	I
3,4-Methylenedioxyamphetamine.	7400	I
4-Methoxyamphetamine	7411	I
Codeine-N-oxide .....	9053	I
Difenoxin .....	9168	I
Heroin .....	9200	I
Morphine-N-oxide .....	9307	I
Norlevorphanol .....	9634	I
Butyryl Fentanyl .....	9822	I
Fentanyl-Related Substance.	9850	I
Methamphetamine .....	1105	II
Lisdexamfetamine .....	1205	II
Phenmetrazine .....	1631	II
Methylphenidate .....	1724	II
Amobarbital .....	2125	II
Pentobarbital .....	2270	II
Secobarbital .....	2315	II
Glutethimide .....	2550	II
Phencyclidine .....	7471	II
4-Anilino-N-phenethyl-4-piperidine (ANPP).	8333	II
Phenylacetone .....	8501	II
Alphaprodine .....	9010	II
Anileridine .....	9020	II
Cocaine .....	9041	II
Dihydrocodeine .....	9120	II
Diphenoxylate .....	9170	II
Levomethorphan .....	9210	II
Levorphanol .....	9220	II
Meperidine .....	9230	II
Dextropropoxyphene, bulk (non-dosage forms).	9273	II
Thebaine .....	9333	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Alfentanil .....	9737	II
Sufentanil .....	9740	II
Tapentadol .....	9780	II

The company plans to import the listed controlled substances for distribution as analytical reference standards to its customers for analytical testing of raw materials. 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.

**Thomas Prevoznik,**

*Deputy Assistant Administrator.*

[FR Doc. 2026-05359 Filed 3-18-26; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1675]

**Bulk Manufacturer of Controlled Substances Application: Sterling Wisconsin, LLC**

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

**ACTION:** Notice of application.

**SUMMARY:** Sterling Wisconsin, LLC 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 18, 2026. Such persons may also file a written request for a hearing on the application on or before May 18, 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 January 27, 2026, Sterling Wisconsin, LLC, W130N10497 Washington Drive, Germantown, Wisconsin 53022-4448, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Lysergic Acid Diethylamide.	7315	I
Marihuana Extract .....	7350	I
Marihuana .....	7360	I
Tetrahydrocannabinols	7370	I
Mescaline .....	7381	I
5-Methoxy-N-N-Dimethyltryptamine.	7431	I
Psilocybin .....	7437	I
Oliceridine .....	9245	II
Thebaine .....	9333	II
Alfentanil .....	9737	II

The company plans to bulk manufacture the listed controlled substances for commercial sale to its customers. In reference to drug codes 7350 (Marihuana Extract), 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

**Thomas Prevoznik,**  
Deputy Assistant Administrator.  
[FR Doc. 2026-05357 Filed 3-18-26; 8:45 am]  
**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1686]

**Importer of Controlled Substances Application: Pisgah Laboratories Inc**

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

**ACTION:** Notice of application.

**SUMMARY:** Pisgah Laboratories Inc 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 20, 2026. Such persons may also file a written request for a hearing on the application on or before April 20, 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 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on January 23, 2026, Pisgah Laboratories Inc, 3222 Old Hendersonville Highway, Pisgah Laboratories Inc, Pisgah Forest, North Carolina 28768, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Phenylacetone .....	8501	II
Ecgonine .....	9180	II
Thebaine .....	9333	II
Tapentadol .....	9780	II

The company plans to manufacture the above-listed controlled substances in bulk for distribution to its customers. 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.

**Thomas Prevoznik,**  
Deputy Assistant Administrator.  
[FR Doc. 2026-05360 Filed 3-18-26; 8:45 am]  
**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

[OMB Number 1122-0001]

**Agency Information Collection Activities; Extension of Previously Approved eCollection eComments Requested; Certification of Compliance With the Statutory Eligibility Requirements of the Violence Against Women Act as Amended, STOP Formula Grant Program**

**AGENCY:** Office on Violence Against Women, Department of Justice.  
**ACTION:** 30-Day notice.

**SUMMARY:** The Department of Justice (DOJ), Office on Violence Against Women, will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995.

**DATES:** Comments are encouraged and will be accepted for 30 days until April 20, 2026.

**FOR FURTHER INFORMATION CONTACT:** If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact: Tiffany Watson, Office on Violence Against Women, at 202-307-6026 or [Tiffany.Watson@usdoj.gov](mailto:Tiffany.Watson@usdoj.gov).

**SUPPLEMENTARY INFORMATION:** The proposed information collection was previously published in the **Federal Register** on January 15, 2026, allowing a 60-day comment period. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Evaluate whether and if so, how the quality, utility, and clarity of the information to be collected can be enhanced; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic,