

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 27, 2025. Such persons may also file a written request for a hearing on the application on or before May 27, 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 30, 2025, 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.

Matthew Strait,
Deputy Assistant Administrator.
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1517]

Importer of Controlled Substances Application: Maridose LLC (I)

AGENCY: Drug Enforcement Administration, Justice.
ACTION: Notice of application.

SUMMARY: Maridose, 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 25, 2025. Such persons may also file a written request for a hearing on the application on or before April 25, 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 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 27, 2025, Maridose, LLC, 74 Orion Street, Unit 7, Brunswick, Maine 04011-5031, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis. In reference to drug codes 7360 (Marihuana) and 7370

(Tetrahydrocannabinols) the company plans to import a synthetic cannabidiol and a synthetic tetrahydrocannabinol. 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. 2025-05065 Filed 3-25-25; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1516]

Importer of Controlled Substances Application: SpecGx LLC

AGENCY: Drug Enforcement Administration, Justice.
ACTION: Notice of application.

SUMMARY: SpecGx 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 25, 2025. Such persons may also file a written request for a hearing on the application on or before April 25, 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 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, 2025, SpecGx LLC, 3600 North 2nd Street, Saint Louis, Missouri 63147-3457, 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
Coca Leaves	9040	II
Thebaine	9333	II
Opium, raw	9600	II
Poppy Straw Concentrate	9670	II
Tapentadol	9780	II

The company plans to import the listed controlled substances for bulk manufacture into Active Pharmaceutical Ingredients for distribution to its customers. In reference to Tapentadol (9780) and Thebaine (9333), the company plans to import intermediate forms of these controlled substances for further manufacturing prior to distribution to its customers. No other activities for these drugs are authorized for this registration. Placement of these codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances.

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. 2025-05060 Filed 3-25-25; 8:45 am]
BILLING CODE P

DEPARTMENT OF JUSTICE

Office of Justice Programs

[OJP (BJA) Docket No. 1835]

Meeting of the Public Safety Officer Medal of Valor Review Board

AGENCY: Bureau of Justice Assistance (BJA), Office of Justice Programs (OJP), Department of Justice (JUSTICE).

ACTION: Notice of meeting.

SUMMARY: This is an announcement of a meeting (via WebEx/conference call-in) of the Public Safety Officer Medal of Valor Review Board to cover a range of issues of importance to the Board, to include but not limited to: Member terms, program administration, marketing, and outreach.

DATES: April 29, 2025, 1:30 p.m. to 2:30 p.m. ET.

ADDRESSES: This meeting will be held virtually using web conferencing technology. The public may hear the proceedings of this virtual meeting/conference call by registering at least seven (7) days in advance with Gregory Joy (contact information below). All emailed requests to register and attend this meeting must include within its Subject line, "MOV Board Meeting April 29, 2025".

FOR FURTHER INFORMATION CONTACT: Gregory Joy, Policy Advisor, Bureau of Justice Assistance, Office of Justice Programs, by telephone at (202) 514-1369, or by email at Gregory.joy@usdoj.gov.

SUPPLEMENTARY INFORMATION: The Public Safety Officer Medal of Valor Review Board carries out those advisory functions specified in 42 U.S.C. 15202. Pursuant to 42 U.S.C. 15201, the President of the United States is authorized to award the Public Safety Officer Medal of Valor, the highest national award for valor by a public safety officer.

This virtual meeting/conference call is open to the public to participate

remotely. For security purposes, members of the public who wish to participate must register at least seven (7) days in advance of the meeting/conference call by contacting Mr. Joy.

Access to the virtual meeting/conference call will not be allowed without prior registration. Please submit any comments or written statements for consideration by the Review Board in writing at least seven (7) days in advance of the meeting date.

Gregory Joy,
Policy Advisor/Designated Federal Officer,
Bureau of Justice Assistance.

[FR Doc. 2025-05052 Filed 3-25-25; 8:45 am]

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NATIONAL LABOR RELATIONS BOARD

Privacy Act of 1974; System of Records

AGENCY: National Labor Relations Board (NLRB).

ACTION: Notice of a new system of records.

SUMMARY: Pursuant to the provisions of the Privacy Act of 1974, the National Labor Relations Board ("NLRB" or "Agency") publishes this notice of a new system of records called "NLRB Freedom of Information Act Records (NLRB-37)." The Agency, elsewhere in the **Federal Register**, is also publishing a notice that it is rescinding two systems of records: NLRB FOIAonline (NLRB-35); and Freedom of Information Act Tracking System (FTS) and Associated Agency Files (NLRB-32). All persons are advised that, in the absence of submitted comments considered by the Agency as warranting modification of the notice as here proposed, it is the intention of the Agency that the notice shall be effective upon expiration of the comment period without further action.

DATES: Written comments on the system's routine uses must be submitted on or before April 25, 2025. The routine uses in this action will become effective on April 25, 2025 unless written comments are received that require a contrary determination.

ADDRESSES: All persons who desire to submit written comments for consideration by the Agency in connection with this proposed notice of the amended system of records shall mail them to the Agency's Senior Agency Official for Privacy, National Labor Relations Board, 1015 Half Street SE, Washington, DC 20570-0001, or submit them electronically to privacy@nlrb.gov. Comments may also be