

| Controlled substance   | Drug code | Schedule |
|--|-----------|----------|
| Gamma Hydroxybutyric Acid .....  | 2010      | I        |
| Methaqualone .....   | 2565      | I        |
| JWH-018 (also known as AM678) (1-Pentyl-3-(1-naphthoyl)indole) .....                       | 7118      | I        |
| JWH-200 (1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl)indole) .....                           | 7200      | I        |
| CP-47,497 (5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol) .....            | 7297      | I        |
| CP-47,497 C8 Homologue (5-(1,1-Dimethyloctyl)-2-[(1R,3S)3-hydroxycyclohexyl]-phenol) ..... | 7298      | I        |
| Lysergic acid diethylamide .....   | 7315      | I        |
| Marihuana .....  | 7360      | I        |
| Tetrahydrocannabinols .....  | 7370      | I        |
| 4-Methyl-2,5-dimethoxyamphetamine .....  | 7395      | I        |
| 3,4-Methylenedioxyamphetamine .....  | 7400      | I        |
| 3,4-Methylenedioxy-N-ethylamphetamine .....  | 7404      | I        |
| 3,4-Methylenedioxymethamphetamine .....  | 7405      | I        |
| Bufotenine .....   | 7433      | I        |
| Psilocybin .....   | 7437      | I        |
| Psilocyn .....   | 7438      | I        |
| Cyprenorphine .....  | 9054      | I        |
| Dihydromorphine .....  | 9145      | I        |
| Heroin .....   | 9200      | I        |
| Normorphine .....  | 9313      | I        |
| Beta-hydroxyfentanyl .....   | 9830      | I        |
| Beta-hydroxy-3-methylfentanyl .....  | 9831      | I        |

The company plans to bulk manufacture the listed controlled substances for the Drug Enforcement Administration-exempted certified reference materials. In-house synthesis gives access to compounds that are difficult to source. In reference to drug codes 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. 2026-04306 Filed 3-3-26; 8:45 am]  
**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1674]

**Importer of Controlled Substances Application: Maridose LLC**

**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 3, 2026. Such persons may also file a written request for a

hearing on the application on or before April 3, 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 January 16, 2026, 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.

**Thomas Prevoznik,**  
Deputy Assistant Administrator.  
[FR Doc. 2026-04307 Filed 3-3-26; 8:45 am]  
**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1671]

**Importer of Controlled Substances Application: Patheon API Inc.**

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

**ACTION:** Notice of application.

**SUMMARY:** Patheon API 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 3, 2026. Such persons may also file a written request for a hearing on the application on or before April 3, 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 9, 2026, Patheon API Inc., 6173 East Old Marion Highway, Florence, South Carolina 29506, 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        |
| Psilocyn .....       | 7438      | I        |

The company plans to import the listed controlled substances as reference standards for research and development as part of API Manufacturing. 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 the Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

**Thomas Prevoznik,**  
Deputy Assistant Administrator.  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1670]

**Importer of Controlled Substances Application: Restek Corporation**

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

**ACTION:** Notice of application.

**SUMMARY:** Restek Corporation 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 3, 2026. Such persons may also file a written request for a hearing on the application on or before April 3, 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, Restek Corporation, 110 Benner Circle, Bellefonte, Pennsylvania 16823-8433, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance  | Drug code | Schedule |
|---|-----------|----------|
| Amineptine .....  | 1219      | I        |
| Mesocarb .....  | 1227      | I        |
| 3-Fluoro-N-methylcathinone (3-FMC) .....  | 1233      | I        |
| Cathinone .....   | 1235      | I        |
| Methcathinone .....   | 1237      | I        |
| 4-Fluoro-N-methylcathinone (4-FMC) .....  | 1238      | I        |
| Para-Methoxymethamphetamine (PMMA), 1-(4-1245 I N methoxyphenyl)-N-methylpropan-2-amine ..... | 1245      | I        |
| Pentadrone (α-methylaminovalerophenone) .....   | 1246      | I        |
| Mephedrone (4-Methyl-N-methylcathinone) .....   | 1248      | I        |
| 4-Methyl-N-ethylcathinone (4-MEC) .....   | 1249      | I        |
| Naphyrone .....   | 1258      | I        |
| N-Ethylamphetamine .....  | 1475      | I        |
| Methiopropamine (N-methyl-1-(thiophen-2-yl)propan-2- 1478 I N amine) .....                    | 1478      | I        |
| N,N-Dimethylamphetamine .....   | 1480      | I        |
| Fenethylamine .....   | 1503      | I        |
| Aminorex .....  | 1585      | I        |
| 4-Methylaminorex (cis isomer) .....   | 1590      | I        |
| 4,4'-Dimethylaminorex .....   | 1595      | I        |
| Gamma Hydroxybutyric Acid .....   | 2010      | I        |