

Signing Authority

This document of the Drug Enforcement Administration was signed on December 15, 2025, by Administrator Terrance C. Cole. That document with the original signature and date is maintained by DEA. For administrative purposes only, and in compliance with requirements of the Office of the Federal Register, the undersigned DEA Federal Register Liaison Officer has been authorized to sign and submit the document in electronic format for publication, as an official document of DEA. This administrative process in no way alters the legal effect of this document upon publication in the **Federal Register**.

Heather Achbach,

Federal Register Liaison Officer, Drug Enforcement Administration.

[FR Doc. 2025–23545 Filed 12–19–25; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE**Drug Enforcement Administration**

[Docket No. DEA–1613]

**Importer of Controlled Substances
Application: National Center for
Natural Products Research**

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: National Center for Natural Products Research 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 January 21, 2026. Such persons may also file a written request for a hearing on the application on or before January 21, 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 September 8, 2025, National Center for Natural Products Research, 806 Hathorn Road, 135 Coy Waller Lab, University, Mississippi 38677, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|-----------------------------|-----------|----------|
| Marihuana | 7360 | I |
| Tetrahydrocannabinols | 7370 | I |

The company plans to acquire new genetic materials with different cannabinoid profiles for research and manufacturing purposes. 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 W. Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2025–23503 Filed 12–19–25; 8:45 am]

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DEPARTMENT OF JUSTICE**Drug Enforcement Administration**

[Docket No. DEA–1636]

**Bulk Manufacturer of Controlled
Substances Application: Navinta LLC**

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Navinta 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 February 20, 2026. Such persons may also file a written request

for a hearing on the application on or before February 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on October 16, 2025,

Navinta LLC, 1499 Lower Ferry Road, Ewing, New Jersey 08618–1414, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|----------------------|-----------|----------|
| Remifentanyl | 9739 | II |

The company plans to bulk manufacture the listed controlled substance for distribution to their customers. No other activity for this drug code is authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2025–23620 Filed 12–19–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1631]

Importer of Controlled Substances Application: Myonex LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Myonex 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 January 21, 2026. Such persons may also file a written request for a hearing on the application on or before January 21, 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 the 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 December 4, 2025, Myonex LLC, 100 Progress Drive, Horsham, Pennsylvania 19044, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|------------------------|-----------|----------|
| Amphetamine | 1100 | II |
| Lisdexamfetamine | 1205 | II |
| Methylphenidate | 1724 | II |
| Nabilone | 7379 | II |
| Oxycodone | 9143 | II |
| Hydromorphone | 9150 | II |
| Hydrocodone | 9193 | II |
| Morphine | 9300 | II |
| Oxymorphone | 9652 | II |
| Fentanyl | 9801 | II |

The company plans to import the listed controlled substances in dosage form for clinical trials, research, and analytical purposes. 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. 2025–23619 Filed 12–19–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1640]

Bulk Manufacturer of Controlled Substances Application: IsoSciences, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: IsoSciences 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 February 20, 2026. Such persons may also file a written request for a hearing on the application on or before February 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on November 18, 2025, IsoSciences, LLC, 340 Mathers Road, Ambler, Pennsylvania 19002–3420, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|----------------------------------|-----------|----------|
| Cathinone | 1235 | I |
| Methcathinone | 1237 | I |
| Lysergic acid diethylamide | 7315 | I |
| Marihuana | 7360 | I |