DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1296]

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 January 23, 2024. Such persons may also file a written request for a hearing on the application on or before January 23, 2024.

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 September 27, 2023, 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
Pentobarbital	2270	II
Levomethorphan	9210	II
Levorphanol	9220	II
Remifentanil	9739	II

The company plans to bulk manufacture the listed controlled substances for validation purposes as part of the Food Administration approval process before distributing to their customers. No other activities for these drug codes are authorized for this registration.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–25991 Filed 11–22–23; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1288]

Bulk Manufacturer of Controlled Substances Application: Organix Chemistry Solutions, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Organix Chemistry Solutions, 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 January 23, 2024. Such persons may also file a written request for a hearing on the application on or before January 23, 2024.

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 September 29, 2023, Organix Chemistry Solutions LLC, 240 Salem Street, Woburn, Massachusetts 01801–2029, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Lysergic acid diethylamide	7315	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Mescaline	7381	I
3,4,5-Trimethoxyamphetamine	7390	I
4-Bromo-2,5-dimethoxyphenethylamine	7392	I
3,4-Methylenedioxyamphetamine	7400	I
3,4-Methylenedioxymethamphetamine	7405	I
5-Methoxy-N-N-dimethyltryptamine	7431	I
Alpha-Methyltryptamine	7432	I
Bufotenine	7433	I
Diethyltryptamine	7434	I
Dimethyltryptamine	7435	I
Psilocybin	7437	I
Psilocyn	7438	I
2-(2,5-Dimethoxy-4-methylphenyl) ethanamine (2C–D)	7508	I
2-(2,5-Dimethoxyphenyl) ethanamine (2C–H)	7517	I
2-(4-Iodo-2,5-dimethoxyphenyl) ethanamine (2C–I)	7518	I
Heroin	9200	I
Morphine	9300	II

The company plans to synthesize the listed controlled substances for distribution to its customers. 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.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–25946 Filed 11–22–23; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1295]

Bulk Manufacturer of Controlled Substances Application: Groff NA Hemplex LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Groff NA Hemplex 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 January 23, 2024. Such persons may also file a written request for a hearing on the application on or before January 23, 2024.

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 September 19, 2023, Groff NA Hemplex LLC., 2218 South Queen Street, York, Pennsylvania 17402, applied to be registered as a bulk manufacturer 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 is federally authorized to conduct cultivation activities in order to bulk manufacture the listed controlled substances for internal use and for sale to federally registered research investigators. No other activities for these drug codes are authorized for this registration.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–25990 Filed 11–22–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1293]

Importer of Controlled Substances Application: Organic Standards Solutions International, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Organic Standards Solutions International, 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 December 26, 2023. Such persons may also file a written request for a hearing on the application on or before December 26, 2023.

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 November 6, 2023, Organic Standards Solutions International, LLC, 7290 Investment Drive, Unit B, North Charleston, South Carolina 29418–8305, 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
Psilocybin	7437	I
Psilocyn	7438	I

The company plans to import the listed controlled substances to produce analytical reference standards for sale and distribution to its customers. Drug codes 7350 (Marihuana Extract) and 7360 (Marihuana) will be used for the manufacture of cannabidiol only. In reference to drug codes 7370 (Tetrahydrocannabinols) the company plans to import a synthetic version of this controlled substance. 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.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–25948 Filed 11–22–23; 8:45 am]

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