

approved finished dosage forms for commercial sale.

**Justin Wood,**  
Acting Deputy Assistant Administrator.  
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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1616]

### Importer of Controlled Substances Application: Irvine Labs, Inc.

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

**ACTION:** Notice of application.

**SUMMARY:** Irvine Labs, 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 December 26, 2025. Such persons may also file a written request for a hearing on the application on or before December 26, 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 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 October 6, 2025, Irvine Labs, Inc., 7305 Murdy Circle, Huntington Beach, California 92647-3533, applied to be registered as an importer 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
Peyote .....	7415	I
Diethyltryptamine .....	7434	I
Dimethyltryptamine .....	7435	I
Psilocybin .....	7437	I
Psilocyn .....	7438	I

The company plans to import bulk substances to support internal research, clinical trials, analytical purposes, and distribution to their customers. In reference to drug codes Marihuana Extract (7350), Marihuana (7360) and Tetrahydrocannabinols (7370), the company plans to import a raw plant material and extracts. 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.

**Justin Wood,**  
Acting Deputy Assistant Administrator.  
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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1618]

### Bulk Manufacturer of Controlled Substances Application: Noramco

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

**ACTION:** Notice of application.

**SUMMARY:** Noramco 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 26, 2026. Such persons may also file a written request for a hearing on the application on or before January 26, 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 September 22, 2025, Noramco, 500 Swedes Landing Road, Wilmington, Delaware 19801-4417, 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
Marihuana .....	7360	I
Tetrahydrocannabinols	7370	I
Dihydromorphine .....	9145	I
Hydromorphone .....	9301	I
Amphetamine .....	1100	II
Lisdexamfetamine .....	1205	II
Methylphenidate .....	1724	II
Nabilone .....	7379	II
Codeine .....	9050	II
Dihydrocodeine .....	9120	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Hydrocodone .....	9193	II
Methadone .....	9250	II
Morphine .....	9300	II
Oripavine .....	9330	II
Thebaine .....	9333	II
Opium extracts .....	9610	II
Opium fluid extract .....	9620	II
Opium tincture .....	9630	II
Opium, powdered .....	9639	II
Opium, granulated .....	9640	II
Opium poppy .....	9650	II
Oxymorphone .....	9652	II
Noroxyphorphone .....	9668	II
Tapentadol .....	9780	II

The company plans to bulk manufacture the listed controlled substances as an Active Pharmaceutical Ingredient for supply 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.

**Justin Wood,**

*Acting Deputy Assistant Administrator.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1617]

#### 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, 2025. Such persons may also file a written request for a hearing on the application on or before December 26, 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 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 October 8, 2025, 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
3,4-Methylenedioxymethamphetamine.	7405	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 code 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.

**Justin Wood,**

*Acting Deputy Assistant Administrator.*

[FR Doc. 2025-21176 Filed 11-25-25; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1607]

#### Bulk Manufacturer of Controlled Substances Application: Siegfried Grafton, Inc.

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

**ACTION:** Notice of application.

**SUMMARY:** Siegfried Grafton, Inc. 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 26, 2026. Such persons may also file a written request for a hearing on the application on or before January 26, 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 September 1, 2025, Siegfried Grafton, Inc., 870 Badger Circle, Grafton, Wisconsin 53024-0000 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
4-Bromo-2,5-dimethoxyphenethylamine.	7392	I
3,4-Methylene dioxyamphetamine.	7400	I
3,4-Methylenedioxymethamphetamine.	7405	I
5-Methoxy-N,N-dimethyltryptamine.	7431	I
Dimethyltryptamine .....	7435	I
Psilocybin .....	7437	I
Psilocyn .....	7438	I
Lisdexamfetamine .....	1205	II
Methylphenidate .....	1724	II
Amobarbital .....	2125	II
Nabilone .....	7379	II
ANPP (4-Anilino-N-phenethyl-4-piperidine).	8333	II
Hydrocodone .....	9193	II