

Controlled substance	Drug code	Schedule
Remifentanyl .....	9739	II

The company plans to import the above listed controlled substance for research purposes.

Dated: November 16, 2018.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2018-25873 Filed 11-26-18; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Registration**

**ACTION:** Notice of registration.

**SUMMARY:** The registrant listed below has applied for and been granted registration by the Drug Enforcement

Administration (DEA) as bulk a manufacturer of various classes of schedule I and II controlled substances.

**SUPPLEMENTARY INFORMATION:** The company listed below applied to be registered as bulk manufacturer of various basic classes of controlled substances. Information on the previously published notice is listed in the table below. No comments or objections were submitted for this notice.

Company	FR Docket	Published
Rhodes Technologies .....	83 FR 40567 .....	August 15, 2018.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of this registrant to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed company.

Dated: November 16, 2018.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2018-25875 Filed 11-26-18; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Application: Cambridge Isotope Laboratories**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before December 27, 2018. Such persons may also file a written request for a hearing on the application on or before December 27, 2018.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also 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.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division ("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on August 30, 2018, Cambridge Isotope Laboratories, 50 Frontage Road, Andover, Massachusetts 01810 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols .....	7370	I
Gamma Hydroxybutyric Acid .....	2010	I
Morphine .....	9300	II

The company plans to import the listed controlled substances for analytical research, testing and clinical trials.

Dated: November 16, 2018.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2018-25869 Filed 11-26-18; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Application: Noramco Inc.**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 28, 2019.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with

respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on July 4, 2018, Noramco Inc., 1550 Olympic Dr. Athens, Georgia 30601 applied to be registered as a bulk manufacturer for the basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Cathinone .....	1235	I
Gamma Hydroxybutyric Acid .....	2010	I
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I
Codeine-N-oxide .....	9053	I
Dihydromorphine .....	9145	I
Hydromorphenol .....	9301	I
Morphine-N-oxide .....	9307	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
Morphine .....	9300	II
Oripavine .....	9330	II
Thebaine .....	9333	II
Opium tincture .....	9630	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Alfentanil .....	9737	II
Sufentanil .....	9740	II
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Fentanyl .....	9801	II

The company plans to manufacture bulk active pharmaceutical ingredients (APIs) and reference standards for distribution to their 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.

Dated: November 16, 2018.

**John J. Martin,**

*Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Registration**

**ACTION:** Notice of registration.

**SUMMARY:** The registrant listed below has applied for and has been granted registration by the Drug Enforcement Administration (DEA) as an importer of schedule I or II controlled substances.

**SUPPLEMENTARY INFORMATION:** The company listed below applied to be registered as an importer of various basic classes of controlled substances. Information on the previously published notice is listed in the table below. No comments or objections were submitted and no requests for a hearing were submitted for this notice.