

Dated: December 3, 2018.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2018–27036 Filed 12–12–18; 8:45 am]

**BILLING CODE 4410–09–P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA–392]

**Bulk Manufacturer of Controlled Substances Application: Eli-Elsohly 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 February 11, 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 November 22, 2017, Eli-Elsohly Laboratories, Mahmoud A. Elsohly Ph.D., 5 Industrial Park Drive, Oxford, Mississippi 38655 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana Extract .....	7350	I
Marihuana .....	7360	I
Tetrahydrocannabinols ..	7370	I
Dihydromorphine .....	9145	I
Amphetamine .....	1100	II
Methamphetamine .....	1105	II

Controlled substance	Drug code	Schedule
Cocaine .....	9041	II
Codeine .....	9050	II
Dihydrocodeine .....	9120	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Ecgonine .....	9180	II
Hydrocodone .....	9193	II
Morphine .....	9300	II
Thebaine .....	9333	II

The company plans to manufacture the listed controlled substances for product development and reference standards. In reference to drug codes 7360 (marihuana) and 7370 (THC), the company plans to isolate these controlled substances from procured 7350 (marihuana extract). In reference to drug code 7360 (marihuana), no cultivation activities are authorized for this registration. No other activities for these drug codes are authorized for this registration.

Dated: December 3, 2018.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2018–27029 Filed 12–12–18; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA–392]

**Bulk Manufacturer of Controlled Substances Application: Cambrex High Point, 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 February 11, 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 16, 2018, Cambrex High Point Inc., 4180 Mendenhall Oaks Parkway, High Point, North Carolina 27265 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substances	Drug code	Schedule
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II

The company plans to manufacture the above listed controlled substances in bulk for distribution to its customers.

Dated: December 3, 2018.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2018–27039 Filed 12–12–18; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA–392]

**Importer 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 14, 2019. Such persons may also file a written request for a hearing on the application on or before January 14, 2019.

**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.