**Authority:** 42 U.S.C. 6213; and 30 CFR 556.511–556.515.

**Walter D. Cruickshank,**

*Acting Director, Bureau of Ocean Energy Management.*

[FR Doc. 2019–24052 Filed 11–4–19; 8:45 am]

**BILLING CODE 4310–MR–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–536]**

### Bulk Manufacturer of Controlled Substances Application: Organix, 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 6, 2020.

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

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on September 9, 2019, Organix, Inc., 240 Salem Street, Woburn, Massachusetts 01801–2029 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Gamma Hydroxybutyric Acid | 2010 | I |
| Lysergic acid diethylamide | 7315 | I |
| Marihuana | 7360 | I |
| Tetrahydrocannabinols | 7370 | I |
| Dimethyltryptamine | 7435 | I |
| Psilocybin | 7437 | I |
| Psilocyn | 7438 | 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 (THC), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: October 18, 2019.

**William T. McDermott,**

*Assistant Administrator.*

[FR Doc. 2019–24107 Filed 11–4–19; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–526]**

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

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturer 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 6, 2020.

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

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on August 6, 2019, Noramco Inc., 500 Swedes Landing Road, Wilmington, Delaware 19801–4417 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Marihuana | 7360 | I |
| Tetrahydrocannabinols | 7370 | I |
| Codeine-N-oxide | 9053 | I |
| Dihydromorphine | 9145 | I |
| Hydromorphinol | 9301 | I |
| Morphine-N-oxide | 9307 | I |
| Amphetamine | 1100 | II |
| Methylphenidate | 1724 | II |
| Nabilone | 7379 | II |
| Phenylacetone | 8501 | 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 extracts | 9610 | II |
| Opium fluid extract | 9620 | II |
| Opium tincture | 9630 | II |
| Opium, powdered | 9639 | II |
| Opium, granulated | 9640 | II |
| Oxymorphone | 9652 | II |
| Noroxymorphone | 9668 | II |
| Tapentadol | 9780 | II |

The company plans to manufacture the listed controlled substances as an Active Pharmaceutical Ingredient (API) for supply to its customers. In reference to drug codes 7360 (marihuana) and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: October 22, 2019.

**William T. McDermott,**

*Assistant Administrator.*

[FR Doc. 2019–24106 Filed 11–4–19; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–530]**

### Importer of Controlled Substances Registration

**ACTION:** Notice of registration.

**SUMMARY:** The registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of schedule I and II controlled substances.

The companies listed below applied to be registered as an importers of various basic classes of schedule I and II controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for a hearing were submitted for these notices.

| Companies | FR docket | Published |
|---|---|---|
| Catalent Pharma Solutions, LLC | 84 FR 36945 | July 30, 2019. |
| Research Triangle Institute | 84 FR 36941 | July 30, 2019. |

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable various basic classes of schedule I and II 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 each of the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying