Controlled substance	Schedule
Bufotenine (7433)	I
Diethyltryptamine (7434)	I
Dimethyltryptamine (7435)	I
Psilocybin (7437)	I
Psilocyn (7438)	I
5-Methoxy-N,N-diisopropyltryptamine (7439)	I
N-Ethyl-1-phenylcyclohexylamine (7455)	I
1-(1-Phenylcyclohexyl)pyrrolidine (7458)	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470)	I
2-(2,5-Dimethoxy-4-ethylphenyl) ethanamine (2C–E) (7509)	I
2-(2,5-Dimethoxyphenyl) ethanamine (2C–H) (7517)	I
2-(4-Iodo-2,5-dimethoxyphenyl) ethanamine (2C–I) (7518)	I
2-(4-Isopropylthio)-2,5-dimethoxyphenyl) ethanamine (2C–T–4) (7532)	I
Dihydromorphine (9145)	I
Heroin (9200)	I
Normorphine (9313)	I
Methamphetamine (1105)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexanecarbonitrile (8603)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Ecgonine (9180)	II
Meperidine intermediate-B (9233)	II
Morphine (9300)	II
Noroxymorphone (9668)	II

The company plans to manufacture high purity drug standards used for analytical applications only in clinical, toxicological, and forensic laboratories and for distribution to its customers.

Dated: January 4, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–00216 Filed 1–8–16; 8:45 am]

BILLING CODE 4410–09–P

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 in accordance with 21 CFR 1301.34(a) on or before February 10, 2016. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before February 10, 2016.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152. Comments and request for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her 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 Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy 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 4, 2015, Noramco, Inc., 500 Swedes Landing Road, Wilmington, Delaware 19801–4417, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Phenylacetone (8501)	II
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780)	II

The company plans to import opium raw (9600) and poppy straw concentrate (9670) to bulk manufacture other controlled substances for distribution to its customers. The company plans to import an intermediate form of tapentadol (9780) to bulk manufacture tapentadol (9780) for distribution to its customers. The company plans to import phenylacetone (8501) in bulk for the manufacture of a controlled substance.

Dated: January 4, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–00209 Filed 1–8–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: IRIX Manufacturing, Inc.

ACTION: Notice of registration.

SUMMARY: IRIX Manufacturing, Inc. applied to be registered as a

manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants IRIX Manufacturing, Inc. registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated August 10, 2015, and published in the **Federal Register** on August 18, 2015, 80 FR 50035, IRIX Manufacturing, Inc., 309 Delaware Street, Building 1106, Greenville, South Carolina 29605 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of IRIX Manufacturing, Inc. to manufacture the 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 above-named company is granted registration as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I

The company plans to manufacture the above-listed controlled substances synthetically as Active Pharmaceutical Ingredients (API) for clinical trials.

Dated: January 4, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–00215 Filed 1–8–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration: Chattem Chemicals Inc.

ACTION: Notice of registration.

SUMMARY: Chattem Chemicals Inc. applied to be registered as an importer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Chattem Chemicals Inc. registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated September 1, 2015, and published in the **Federal Register** on September 9, 2015, 80 FR 54326, Chattem Chemicals Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409 applied to be registered as an importer of certain basic classes of controlled substances. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007). No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Chattem Chemicals Inc. to import the 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above-named company is granted registration as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Methamphetamine (1105)	II
4-Anilino-N-phenethyl-4-piperidine (ANPP) (8333).	II
Phenylacetone (8501)	II
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780)	II

The company plans to import the listed controlled substances to bulk manufacture other controlled substances for distribution to its customers. The company plans to import an intermediate form of tapentadol (9780), to bulk manufacture tapentadol (9780) for distribution to its customers. The company plans to import phenylacetone (8501) in bulk for the manufacture of a controlled substance.

Dated: January 4, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–00218 Filed 1–8–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: AMPAC Fine Chemicals LLC

ACTION: Notice of registration.

SUMMARY: AMPAC Fine Chemicals LLC applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants AMPAC Fine Chemicals LLC registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated June 25, 2015, and published in the **Federal Register** on July 6, 2015, 80 FR 38467, AMPAC Fine Chemicals LLC, Highway 50 and Hazel Avenue, Building 05001, Rancho Cordova, California 95670 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of AMPAC Fine Chemicals LLC to manufacture the 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 above-named company is granted registration as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Methylphenidate (1724)	II
Thebaine (9333)	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780)	II

The company is a contract manufacturer. In reference to Poppy Straw Concentrate the company will manufacture thebaine intermediates for sale to its customers for further manufacture. No other activity for this drug code is authorized for this registration.