

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Manufacturer of Controlled Substances Registration: Chattem Chemicals, Inc.**

**ACTION:** Notice of registration.

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

**SUPPLEMENTARY INFORMATION:** By notice dated October 13, 2015, and published in the **Federal Register** on October 21, 2015, 80 FR 63835, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409 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 Chattem Chemicals, 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
Gamma Hydroxybutyric Acid (2010)	I
4-Methoxyamphetamine (7411) ...	I
Dihydromorphine (9145) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Lisdexamfetamine (1205) .....	II
Methylphenidate (1724) .....	II
Pentobarbital (2270) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Meperidine intermediate-A (9232)	II
Meperidine intermediate-B (9233)	II

Controlled substance	Schedule
Meperidine intermediate-C (9234)	II
Methadone (9250) .....	II
Methadone intermediate (9254) ...	II
Morphine (9300) .....	II
Oripavine (9330) .....	II
Thebaine (9333) .....	II
Opium tincture (9630) .....	II
Opium, powdered (9639) .....	II
Opium, granulated (9640) .....	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Remifentanil (9739) .....	II
Sufentanil (9740) .....	II
Tapentadol (9780) .....	II
Fentanyl (9801) .....	II

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

Dated: March 14, 2016.  
**Louis J. Milione,**  
*Deputy Assistant Administrator.*  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Registration: Noramco, Inc.**

**ACTION:** Notice of registration.

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

**SUPPLEMENTARY INFORMATION:** By notice dated August 21, 2015, and published in the **Federal Register** on August 31, 2015, 80 FR 52510, Noramco, Inc., 1440 Olympic Drive, Athens, Georgia 30601 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 Noramco, 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
Phenylacetone (8501) .....	II
Thebaine (9333) .....	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780) .....	II

The company plans to import thebaine (9333) analytical reference standards for distribution to its customers. The company plans to import an intermediate form of tapentadol (9780) to bulk manufacture tapentadol for distribution to its customers. The company plans to import phenylacetone (8501) and poppy straw concentrate (9670) to manufacture other controlled substances.

Dated: March 14, 2016.  
**Louis J. Milione,**  
*Deputy Assistant Administrator.*  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Manufacturer of Controlled Substances Registration: Cerilliant Corporation**

**ACTION:** Notice of registration.

**SUMMARY:** Cerilliant Corporation applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Cerilliant Corporation registration as a manufacturer of those controlled substances.

**SUPPLEMENTARY INFORMATION:** By notice dated September 21, 2015, and published in the **Federal Register** on September 30, 2015, 80 FR 58788, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665-2402 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments