

Cambrex Charles City, Inc. to import the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Cambrex Charles City, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with State and local laws, and a review of 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 basic class of controlled substance listed.

Dated: May 14, 2007.

**Joseph T. Rannazzisi,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated October 25, 2006, and published in the **Federal Register** on November 1, 2006, (71 FR 64298), Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Sufentanil (9740), a basic class of controlled substance listed in schedule II.

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

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cambrex Charles City, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Cambrex Charles City, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with State and local laws, and a review of the company's background and history.

Therefore, pursuant to 21 U.S.C. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: May 14, 2007.

**Joseph T. Rannazzisi,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated January 16, 2007, and published in the **Federal Register** on January 23, 2007, (72 FR 2907), Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedule I and II:

Drug	Schedule
Tetrahydrocannabinols (7370) .....	I
Dihydromorphine (9145) .....	I
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II
Remifentanil (9739) .....	II

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

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cedarburg Pharmaceuticals, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Cedarburg Pharmaceuticals, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with State and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of

the basic classes of controlled substances listed.

Dated: May 14, 2007.

**Joseph T. Rannazzisi,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated November 21, 2006, and published in the **Federal Register** on December 1, 2006, (71 FR 69589-69590), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal and by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedule I and II:

Drug	Schedule
4-Methoxyamphetamine (7411) ...	I
Dihydromorphine (9145) .....	I
Difenoxin (9168) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	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
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

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

By letter dated February 15, 2007, Chattem Chemicals has withdrawn their request for N-Ethylamphetamine (1475), Secobarbital (2315), 2,5-Dimethoxyamphetamine (7396), Diphenoxylate (9170), Opium Extracts (9610), Opium Fluid Extract (9620), Opium Tincture (9630), Opium Powdered (9639), and Opium Granulated (9640).

One comment has been received; this comment has been noted and considered by DEA.