

DEPARTMENT OF JUSTICE**Drug Enforcement Administration****Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated September 28, 2011, and published in the **Federal Register** on October 7, 2011, 76 FR 62449, Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (GHB) (2010)	I
Amphetamine (1100)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
4-Anilino-N-phenethyl-4-piperidine (8333)	II
Phenylacetone (8501)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Opium raw (9600)	II
Oxymorphone (9652)	II
Poppy Straw Concentrate (9670)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company will manufacture the listed controlled substances in bulk for sale 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 classes of controlled substances 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 USC § 823(a), 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: January 27, 2012.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2012-2591 Filed 2-3-12; 8:45 am]

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DEPARTMENT OF JUSTICE**Drug Enforcement Administration****Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated September 28, 2011, and published in the **Federal Register** on October 7, 2011, 76 FR 62450, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to manufacture a radioactive product to diagnose Parkinson's disease, and to manufacture a bulk investigational new drug (IND) for clinical trials.

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 GE Healthcare to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated GE Healthcare 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: January 26, 2012.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2012-2586 Filed 2-3-12; 8:45 am]

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DEPARTMENT OF JUSTICE**Drug Enforcement Administration****Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated June 23, 2011, and published in the **Federal Register** on July 5, 2011, 76 FR 39127, Johnson Matthey Pharmaceutical Materials, Inc., Pharmaceuticals Service, 25 Patton Road, Devens, Massachusetts 01434, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Remifentanyl (9739), the basic class of controlled substance in schedule II.

The company plans to utilize this facility to manufacture small quantities of the listed controlled substance in bulk and to conduct analytical testing in support of the company's primary manufacturing facility in West Deptford, New Jersey. The controlled substances manufactured in bulk at this facility will be distributed to the company's 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 Johnson Matthey Pharmaceutical Materials, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Johnson Matthey Pharmaceutical Materials, 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: January 27, 2012.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2012-2568 Filed 2-3-12; 8:45 am]

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DEPARTMENT OF JUSTICE**Parole Commission****Notice of Sunshine Act Meeting**

TIME AND DATE: 11:30 a.m., Thursday, February 9, 2012.