

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; GE Healthcare

By Notice dated November 2, 2013, and published in the Federal Register on November 19, 2013, 78 FR 69447, 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 an importer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to import small quantities of ioflupane, in the form of three separate analogues of Cocaine, that will be used for the support and manufacture of DaTSCAN (ioflupane I-123) injections for distribution as a radioactive diagnostic imaging agent utilized in the diagnosis of Parkinson's disease.

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of GE Healthcare 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. The 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. 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: February 19, 2014.

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

[FR Doc. 2014-05480 Filed 3-12-14; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Siegfried USA, LLC

By Notice dated December 23, 2013, and published in the Federal Register on January 10, 2014, 79 FR 1887, Siegfried USA, LLC., 33 Industrial Park Road, Pennsville, New Jersey 08070, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Table with 2 columns: Drug, Schedule. Rows: Opium, raw (9600) II; Poppy Straw Concentrate (9670) II

The company plans to import the listed controlled substances to bulk manufacture APIs for distribution to its customer.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

The DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Siegfried USA, LLC., 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 has investigated Siegfried USA, LLC., 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 classes of controlled substances listed.

Dated: February 19, 2014.

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

[FR Doc. 2014-05497 Filed 3-12-14; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Johnson Matthey, Inc.

By Notice dated November 4, 2013, and published in the Federal Register on November 18, 2013, 78 FR 69130, Johnson Matthey, Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Table with 2 columns: Drug, Schedule. Rows: Coca Leaves (9040) II; Thebaine (9333) II; Opium, raw (9600) II; Noroxymorphone (9668) II; Poppy Straw Concentrate (9670) II

The company plans to import the listed controlled substances as raw materials, to be used in the manufacture of bulk controlled substances, for distribution to its customers.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417, (January 25, 2007).

In reference to the non-narcotic raw material, no comments or objections have been received. The company plans to import gram amounts to be used as reference standards for distribution to its customers.

The DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Johnson Matthey, 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 has investigated Johnson Matthey, 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 classes of controlled substances listed.

Dated: February 19, 2014.

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

[FR Doc. 2014-05488 Filed 3-12-14; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

**Manufacturer of Controlled
Substances; Notice of Application;
Cayman Chemical Company**

Pursuant to 21 CFR 1301.33(a), this is notice that on December 6, 2013, Cayman Chemical Company, 1180 East Ellsworth Road, Ann Arbor, Michigan 48108, made application by correspondence to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
2-(4-bromo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) (25B-NBOMe) (7536).	I
2-(4-chloro-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) (25C-NBOMe) (7537).	I
2-(4-iodo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) (25I-NBOMe) (7538).	I

The company plans to manufacture the listed controlled substances, and distribute those substances to its research and forensics customers for drug testing and analysis.

Any other such applicant, and any person who is presently registered with the DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODW), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than May 12, 2014.

Dated: Signed February 19, 2014.

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

[FR Doc. 2014-05493 Filed 3-12-14; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

**Manufacturer of Controlled
Substances; Notice of Registration;
Johnson Matthey, Inc.**

By Notice dated November 5, 2013, and published in the **Federal Register** on November 18, 2013, 78 FR 69132, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742, 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 (2010).	I
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Propiram (9649)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II
Morphine (9300)	II
Thebaine (9333)	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 sale to its customers.

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Johnson Matthey, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. The DEA has investigated, Johnson Matthey, 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 classes of controlled substances listed.

Dated: February 19, 2014.

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;
Cedarburg Pharmaceuticals, Inc.**

By Notice dated October 16, 2013, and published in the **Federal Register** on October 25, 2013, 78 FR 64017, 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 following basic classes of controlled substances:

Drug	Schedule
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Remifentanil (9739)	II
Fentanyl (9801)	II

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

No comments or objections have been received. The 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 classes of controlled substances is consistent with the public interest at this time. The 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(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.