DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Mylan Technologies, Inc.

By Notice dated January 15, 2013, and published in the **Federal Register** on January 29, 2013, 78 FR 6131, Mylan Technologies, Inc., 110 Lake Street, Saint Albans, Vermont 05478, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Methylphenidate (1724)	II
Fentanyl (9801)	II

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to afford the company the opportunity to export domestically-manufactured FDF to foreign markets.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Mylan Pharmaceuticals, 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. DEA has investigated Mylan 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. 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: May 24, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013–13181 Filed 6–3–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Caraco Pharmaceutical Laboratories, LTD

By Notice dated February 8, 2013, and published in the **Federal Register** on February 21, 2013, 78 FR 12101, Caraco Pharmaceutical Laboratories, Ltd., 270 Prospect Plains Road, Cranbury, New Jersey 08512, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Fentanyl (9801), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance in finished dosage form for clinical trials, and research.

The import of the above listed basic class of controlled substance is granted only for analytical testing and clinical trials. This authorization does not extend to the import of finished FDA approved or non-approved dosage form for commercial distribution in the United States.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Caraco Pharmaceutical Laboratories, Ltd., 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. DEA has investigated Caraco Pharmaceutical Laboratories, Ltd., 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 24, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013–13183 Filed 6–3–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Agilent Technologies

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 27, 2013, Agilent Technologies, 25200 Commercentre Drive, Lake Forest, California 92630–8810, 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 II:

Drug	Schedule
1-Piperidinocyclohexane-carbonitrile (8603)	II
Benzoylecgonine (9180)	II

The company plans to manufacture small quantities of the listed controlled substances for use in diagnostic products.

Any other such applicant, and any person who is presently registered with 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 (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 5, 2013.

Dated: May 24, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013–13219 Filed 6–3–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Penick Corporation

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 23, 2013, Penick Corporation, 33 Industrial Park Road, Pennsville, New Jersey 08070, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Opium Tincture (9630), a basic class of

controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance, as bulk intermediates for distribution to its customers.

Any other such applicant, and any person who is presently registered with 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 (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 5, 2013.

Dated: May 24, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013–13221 Filed 6–3–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application Sigma Aldrich Research Biochemicals, Inc.

Pursuant to 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 7, 2013, Sigma Aldrich Research Biochemicals, Inc., 1–3 Strathmore Road, Natick, Massachusetts 01760–2447, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Mephedrone (4-Methyl-N-methylcathinone) (1248).	I
MDPV (3,4-Methylenedioxypyrovalerone) (7535).	I
Methylone (3,4-Methylenedioxy-N-methylcathinone) (7540).	I
Sufentanil (9740)	II

The company plans to manufacture reference standards.

Any other such applicant, and any person who is presently registered with 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 (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 5, 2013.

Dated: May 24, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013–13217 Filed 6–3–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Cerilliant Corporation

By Notice dated January 14, 2013, and published in the **Federal Register** on January 25, 2013, 78 FR 5499, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665–2402, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
JWH–250 (6250)	I
SR–18 also known as RCS–8 (7008).	I
JWH–019 (7019)	I
JWH–081 (7081)	I
SR–19 also known as RCS–4 (7104).	I
JWH–122 (7122)	I
AM–2201 (7201)	I
JWH–203 (7203)	I
2C–T–2 (7385)	I
JWH–398 (7398)	I
N-Ethyl-1-phenylcyclohexylamine (7455).	I
2C–D (7508)	I
2C–E (7509)	I
2C–H (7517)	I
2C–I (7518)	I
2C–C (7519)	I
2C–N (7521)	I
2C–P (7524)	I
2C–T–4 (7532)	I
AM–694 (7694)	I
Metazocine (9240)	II

The company plans to manufacture the listed controlled substances for distribution to their research and forensic customers conducting drug testing and analysis.

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 Cerilliant Corporation to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cerilliant Corporation 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: May 24, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013–13225 Filed 6–3–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; GE Healthcare

By Notice dated February 8, 2013 and published in the **Federal Register** on February 21, 2013, 78 FR 12103, 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