

Drug	Schedule
Amphetamine (1100) .....	II
Lisdexamfetamine (1205) .....	II
Methylphenidate (1724) .....	II
Pentobarbital (2270) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Levorphanol (9220) .....	II
Morphine (9300) .....	II
Oxymorphone (9652) .....	II
Remifentanyl (9739) .....	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 allow the company 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: December 31, 2013.

**Joseph T. Rannazzisi,**

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

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Registration; Fisher Clinical Services, Inc.

By Notice dated September 16, 2013, and published in the **Federal Register**

on September 25, 2013, 78 FR 59064, Fisher Clinical Services, Inc., 7554 Schantz Road, Allentown, Pennsylvania 18106, 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
Methyphenidate (1724) .....	II
Levorphanol (9220) .....	II
Noroxymorphone (9668) .....	II
Tapentadol (9780) .....	II

The company plans to import the listed controlled substances for clinical trials, analytical research and testing.

The import of the above listed basic classes of controlled substances will be granted only for analytical testing and clinical trials. This authorization does not extend to the import of a 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 Fisher Clinical Services, 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 Fisher Clinical Services, 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: December 31, 2013.

**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 Application; Cayman Chemical Company

Pursuant to 21 CFR 1301.33(a), this is notice that on June 27, 2013, Cayman Chemical Company, 1180 East Ellsworth Road, Ann Arbor, Michigan 48108, 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
Cathinone (1235) .....	I
Methcathinone (1237) .....	I
4-Methyl-N-methylcathinone (1248) .....	I
N-Ethylamphetamine (1475) .....	I
N,N-Dimethylamphetamine (1480) .....	I
Aminorex (1585) .....	I
4-Methylaminorex (cis isomer) (1590) .....	I
Gamma Hydroxybutyric Acid (2010) .....	I
JWH-250 (6250) .....	I
SR-18 also known as RCS-8 (7008) .....	I
XLR11 (7011) .....	I
JWH-019 (7019) .....	I
AKB48 (7048) .....	I
JWH-081 (7081) .....	I
SR-19 also known as RCS-4 (7104) .....	I
1-Pentyl-3-(1-naphthoyl)indole (7118) .....	I
JWH-122 (7122) .....	I
UR-144 (7144) .....	I
1-Butyl-3-(1-naphthoyl)indole (7173) .....	I
1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl) indole (7200) .....	I
AM-2201 (7201) .....	I
JWH-203 (7203) .....	I
Alpha-ethyltryptamine (7249) .....	I
5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol (7297) .....	I
5-(1,1-Dimethyloctyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol (7298) .....	I
Lysergic acid diethylamide (7315) .....	I
2,5-Dimethoxy-4-(n-propylthiophenethylamine (7348) .....	I
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	I
Mescaline (7381) .....	I
2C-T-2 (7385) .....	I
3,4,5-Trimethoxyamphetamine (7390) .....	I
4-Bromo-2,5-dimethoxyamphetamine (7391) .....	I
4-Bromo-2,5-dimethoxyphenethylamine (7392) .....	I
4-Methyl-2,5-dimethoxyamphetamine (7395) .....	I