

that on September 12, 2011, Mallinckrodt LLC., 3600 North Second Street, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Drug	Schedule
Methylphenidate (1724) .....	II
Phenylacetone (8501) .....	II
Coca Leaves (9040) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II
Opium, raw (9600) .....	II
Poppy Straw Concentrate (9670) .....	II
Fentanyl (9801) .....	II

The company plans to import the listed controlled substances to manufacture bulk controlled substances for distribution to its customers, and for research and analytical standards.

No comments, objections, or requests for any hearings will be accepted on any application for registration or re-registration to import crude opium, poppy straw, poppy straw concentrate, and coca leaves. As explained in the Correction to Notice of Application pertaining to 72 FR 3417 (2007), comments and requests for hearings on applications to import narcotic raw material are not appropriate.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule I or II, which fall under the authority of section 1002(a)(2)(B) of the Act 21 U.S.C. 952 (a)(2)(B) may, in the circumstances set forth in 21 U.S.C. 958(i), file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

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 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than March 1, 2012.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f).

As noted in a previous notice published in the **Federal Register** on September 23, 1975, 40 FR 43745–46, all applicants for registration to import a basic class of any controlled substance

in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: January 23, 2012.

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

[FR Doc. 2012–1980 Filed 1–30–12; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Registration

By Notice dated June 1, 2011, and published in the **Federal Register** on June 9, 2011, 76 FR 33784, Alltech Associates, Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, 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
Gamma Hydroxybutyric Acid (2010) .....	I
Heroin (9200) .....	I
Lysergic acid diethylamide (7315) .....	I
Cocaine (9041) .....	II
Codeine (9050) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II

The company plans to import these controlled substances for the manufacture of reference standards.

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 Alltech Associates, 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 Alltech Associates, 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: January 23, 2012.

**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 June 23, 2011, and published in the **Federal Register** on July 5, 2011, 76 FR 39127, 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
N-Ethylamphetamine (1475) .....	I
N,N-Dimethylamphetamine (1480) .....	I
Aminorex (1585) .....	I
4-Methylaminorex (cis isomer) (1590) .....	I
1-Pentyl-3-(1-naphthoyl)indole (7118) .....	I
1-Butyl-3-(1-naphthoyl)indole (7173) .....	I
1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl) Indole (7200) .....	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-propylthiophenethyl)amine (7348) .....	I
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	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
2,5-Dimethoxyamphetamine (7396) .....	I

Drug	Schedule
2,5-Dimethoxy-4-ethylamphetamine (7399).	I
3,4-Methylenedioxyamphetamine (7400).	I
5-Methoxy-3,4-methylenedioxyamphetamine (7401).	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymethamphetamine (7405).	I
4-Methoxyamphetamine (7411) ...	I
5-Methoxy-N-N-dimethyltryptamine (7431).	I
Alpha-methyltryptamine (7432) ....	I
Diethyltryptamine (7434) .....	I
Dimethyltryptamine (7435) .....	I
5-Methoxy-N,N-diisopropyltryptamine (7439).	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Lisdexamfetamine (1205) .....	II

The company plans to manufacture small quantities of marijuana derivatives for research purposes. In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol. In reference to drug code 7370 (Tetrahydrocannabinols), the company will manufacture a synthetic THC. No other activity for this drug code is authorized for this registration.

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 Cayman Chemical Company to manufacture the listed basic classed of controlled substances is consistent with the public interest at this time. DEA has investigated Cayman Chemical Company 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: January 23, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-1977 Filed 1-30-12; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 7, 2011, and published in the **Federal Register** on June 16, 2011, 76 FR 35243, Archimica, Inc., 2460 W. Bennett Street, Springfield, Missouri 65807-1229, 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
Amphetamine (1100) .....	II
Lisdexamfetamine (1205) .....	II
Methylphenidate (1724) .....	II
Phenylacetone (8501) .....	II
Methadone Intermediate (9254) ...	II
Tapentadol (9780) .....	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers for Amphetamine (1100).

The company plans to acquire the listed controlled substances in bulk from a domestic source in order to manufacture other 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 Archimica, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Archimica, 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: January 23, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-1981 Filed 1-30-12; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Federal Bureau of Investigation

[OMB Number 1110-0004]

#### Agency Information Collection Activities; Proposed Collection, Comments Requested: Extension of a Currently Approved Collection; Number of Full-Time Law Enforcement Employees as of October 31

**ACTION:** 30-day Notice of Information Collection Under Review.

The Department of Justice, Federal Bureau of Investigation, Criminal Justice Information Services Division (CJIS) will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and clearance in accordance with established review procedures of the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register** Volume 76, Number 228, page 72977, on November 28, 2011, allowing for a 60 day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until March 1, 2012. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to Mr. Gregory E. Scarbro, Unit Chief, Federal Bureau of Investigation, CJIS Division, Module E-3, 1000 Custer Hollow Road, Clarksburg, West Virginia 26306; facsimile (304) 625-3566.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Comments should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and