**Summary of Information Collection**

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Notification of Change of Mailing or Premise Address.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: None. Bureau of Alcohol, Tobacco, Firearms and Explosives.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Not-for-profit-institutions. Other: Business or other for-profit.

**Need for Collection**

Licensees and permittees whose mailing address will change must notify the Chief, Federal Explosives Licensing Center, at least 10 days before the change. The information is used by ATF to identify correct locations of storage of explosives licensees/permittees and location of storage of explosive materials for purposes of inspection, as well as to notify permittee/licensees of any change in regulations or laws that may affect their business activities.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that 1,000 respondents will take 10 minutes to respond via letter to the Federal Explosives Licensing Center.

(6) *An estimate of the total public burden (in hours) associated with the collection:* There are an estimated 170 annual total burden hours associated with this collection.

*If additional information is required contact:* Jerri Murray, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, Department of Justice, Two Constitution Square, Room 2E–508, 145 N Street NE., Washington, DC 20530.

**Jerri Murray,**

*Department Clearance Officer, PRA, U.S. Department of Justice.*

[FR Doc. 2012–12346 Filed 5–21–12; 8:45 am]

**BILLING CODE 4410–FY–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Importer of Controlled Substances; Notice Of Registration; Mylan Pharmaceuticals, Inc.

By Notice dated February 23, 2012, and published in the **Federal Register** on March 1, 2012, 77 FR 12620, Mylan Pharmaceuticals, Inc., 781 Chestnut Ridge Road, Morgantown, West Virginia 26505, made application by letter to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

| Drug | Schedule |
|---|---|
| Amphetamine (1100) | II |
| Methadone (9250) | II |
| Morphine (9300) | 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: May 15, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012–12382 Filed 5–21–12; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Application; Lin Zhi International Inc.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 19, 2012, Lin Zhi International Inc., 670 Almanor Avenue, Sunnyvale, California 94085, 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 |
|---|---|
| Tetrahydrocannabinols (7370) | I |
| 3,4–Methylenedioxymethamphetamine (7405). | I |
| Cocaine (9041) | II |
| Oxycodone (9143) | II |
| Hydrocodone (9193) | II |
| Methadone (9250) | II |
| Dextropropoxyphene, bulk (non-dosage forms) (9273). | II |
| Morphine (9300) | II |

The company plans to manufacture the listed controlled substances as bulk reagents for use in drug abuse testing.

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 July 23, 2012.

Dated: May 15, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012–12326 Filed 5–21–12; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice Of Application; Research Triangle Institute

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 12, 2012, Research Triangle Institute, Hermann Building, East Institute Drive, P.O. Box 12194, Research Triangle, North Carolina 27709, 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 |
|---|---|
| Marihuana (7360) | I |
| Cocaine (9041) | II |

The Institute will manufacture marihuana, and cocaine derivatives for use by their customers in analytical kits, reagents, and reference standards as directed by the National Institute on Drug Abuse.

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 July 23, 2012.

Dated: May 15, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012–12381 Filed 5–21–12; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances, Notice of Application, Alltech Associates, Inc.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 19, 2012, Alltech Associates Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, 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 |
|---|---|
| Methcathinone (1237) | I |
| N-Ethylamphetamine (1475) | I |
| N,N-Dimethylamphetamine (1480) | I |
| 4-Methylaminorex (cis isomer) (1590). | I |
| Alpha-ethyltryptamine (7249) | I |
| Lysergic acid diethylamide (7315) | I |
| 2,5-Dimethoxy-4-(n)-propylthiophenethylamine (7348). | I |
| Tetrahydrocannabinols (7370) | I |
| Mescaline (7381) | I |
| 4-Bromo-2,5-dimethoxy-amphetamine (7391). | I |
| 4-Bromo-2,5-dimethoxyphenethylamine (7392). | I |
| 4-Methyl-2,5-dimethoxy-amphetamine (7395). | I |
| 2,5-Dimethoxyamphetamine (7396). | I |
| 2,5-Dimethoxy-4-ethylamphetamine (7399). | I |
| 3,4-Methylenedioxyamphetamine (7400). | I |
| N-Hydroxy-3,4-methylenedioxyamphetamine (7402). | I |
| 3,4-Methylenedioxy-N-ethylamphetamine (7404). | I |
| 3,4-Methylenedioxymethamphetamine (7405). | I |
| 4-Methoxyamphetamine (7411) | I |
| Alpha-methyltryptamine (7432) | I |
| Bufotenine (7433) | I |
| Diethyltryptamine (7434) | I |
| Dimethyltryptamine (7435) | I |
| Psilocybin (7437) | I |
| Psilocyn (7438) | I |
| 5-Methoxy-N,N-diisopropyltryptamine (7439). | I |
| N-Ethyl-1-phenylcyclohexylamine (7455). | I |
| 1-(1-Phenylcyclohexyl)pyrrolidine (7458). | I |
| 1-[1-(2-Thienyl)-cyclohexyl]piperidine (7470). | I |
| Dihydromorphine (9145) | I |
| Heroin (9200) | I |
| Normorphine (9313) | I |
| Methamphetamine (1105) | II |
| 1-phenylcyclohexylamine (7460) II | II |
| Phencyclidine (7471). | |
| Phenylacetone (8501) | II |
| 1-piperidinocyclohexanecarbonitrile (8603). | II |
| Cocaine (9041) | II |
| Codeine (9050) | II |
| Dihydrocodeine (9120) VII. | |
| Ecgonine (9180) | II |
| Meperidine intermediate-B (9233) | II |
| Noroxymorphone (9668) VII. | |

The company plans to manufacture high purity drug standards used for analytical applications only in clinical, toxicological, and forensic laboratories.

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 July 23, 2012.

Dated: May 15, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012–12379 Filed 5–21–12; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF LABOR

### Approval of Information Collection Requirements; Comment Request

**AGENCY:** Office of Federal Contract Compliance Programs, Labor.

**ACTION:** Notice.

**SUMMARY:** The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a pre-clearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA). 44 U.S.C. 3506(c)(2)(A). This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. Currently, the Office of Federal Contract Compliance Programs is soliciting comments on its