

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, 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 26, 2010.

**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 October 20, 2009, and published in the **Federal Register** on October 28, 2009, (74 FR 55588), Aldrich Chemical Company, Inc., DBA Isotec, 3858 Benner Road, Miamisburg, Ohio 45342-4304, 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 schedules I and II:

| Drug  | Schedule |
|---|----------|
| Gamma Hydroxybutyric Acid (2010)                          | I        |
| Methaqualone (2565)                                       | I        |
| l-bogaine (7260)  | I        |
| Tetrahydrocannabinols (7370)                              | I        |
| 2,5-Dimethoxyamphetamine (7396)                           | I        |
| Psilocyn (7438)   | I        |
| Normorphine (9313)  | I        |
| Acetylmethadol (9601)                                     | I        |
| Alphacetylmethadol except levo-alpha-cetylmethadol (9603) | I        |
| Normethadone (9635)                                       | I        |
| Norpipanone (9636)  | I        |
| 3-Methylfentanyl (9813)                                   | I        |
| Amphetamine (1100)  | II       |
| Methamphetamine (1105)                                    | II       |
| Methylphenidate (1724)                                    | II       |
| Amobarbital (2125)  | II       |
| Pentobarbital (2270)                                      | II       |
| Secobarbital (2315)                                       | II       |
| 1-Phenylcyclohexylamine (7460)                            | II       |
| Phencyclidine (7471)                                      | II       |
| Phenylacetone (8501)                                      | II       |
| 1-Piperidinocyclohexanecarbonitrile (8603)                | II       |
| Cocaine (9041)  | II       |

| Drug  | Schedule |
|---|----------|
| Codeine (9050)                                      | II       |
| Oxycodone (9143)                                    | II       |
| Hydromorphone (9150)                                | II       |
| Benzoylcegonine (9180)                              | II       |
| Ethylmorphine (9190)                                | II       |
| Hydrocodone (9193)                                  | II       |
| Isomethadone (9226)                                 | II       |
| Meperidine (9230)                                   | II       |
| Meperidine intermediate-A (9232)                    | II       |
| Meperidine intermediate-B (9233)                    | II       |
| Methadone (9250)                                    | II       |
| Methadone intermediate (9254)                       | II       |
| Dextropropoxyphene, bulk, (non-dosage forms) (9273) | II       |
| Morphine (9300)                                     | II       |
| Thebaine (9333)                                     | II       |
| Levo-alpha-cetylmethadol (9648)                     | II       |
| Oxymorphone (9652)                                  | II       |

The company plans to manufacture small quantities of the listed controlled substances to produce isotope labeled standards for 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 Aldrich Chemical Company Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Aldrich Chemical Company 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, 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 26, 2010.

**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 October 16, 2009, and published in the **Federal Register** on October 28, 2009, (74 FR 55586), Lin Zhi International Inc., 687 North Pastoria Avenue, Sunnyvale, California 94085, 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 schedules I and II:

| Drug   | Schedule |
|--|----------|
| Tetrahydrocannabinols (7370)                       | I        |
| 3,4-Methylenedioxymethamphetamine (MDMA) (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.

No comments or objections have been received. DEA has considered the factors in 21 USC 823(a) and determined that the registration of Lin Zhi International Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Lin Zhi International 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 USC 823, 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 26, 2010.

**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 September 17, 2009, and published in the **Federal Register** on September 25, 2009, (74 FR 49020), Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665-2402, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes