

| Drug | Schedule |
|--|----------|
| 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 |
| Codeine (9050) | II |
| Dihydrocodeine (9120) | II |
| Oxycodone (9143) | II |
| Hydromorphone (9150) | II |
| Benzoyllecgonine (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-dosageforms) (9273) | II |
| Morphine (9300) | II |
| Thebaine (9333) | II |
| Levo-alphaacetylmethadol (9648) | II |
| Oxymorphone (9652) | II |
| Fentanyl (9801) | 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: November 5, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-22475 Filed 11-15-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II and prior to issuing a registration under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on October 5, 2007, JFC Technologies LLC., 100 West Main Street, P.O. Box 669, Bound Brook, New Jersey 08805, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Meperidine intermediate-B (9233), a basic class of controlled substance listed in schedule II.

The company plans to import the basic class of controlled substance for the production of controlled substances for clinical trials, research, analytical purposes, and distribution to its customers.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class of controlled substance

may 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 comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, VA. 22152; and must be filed no later than December 17, 2007.

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 substances 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: November 6, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-22488 Filed 11-15-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 24, 2007, JFC Technologies, LLC., 100 W. Main Street, Bound Brook, New Jersey 08805, 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 |
|----------------------------|----------|
| Diphenoxylate (9170) | II |
| Hydrocodone (9193) | II |

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance 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 being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, VA 22152; and must be filed no later than January 15, 2008.

Dated: November 5, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-22520 Filed 11-15-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 26, 2007, and published in the **Federal Register** on July 3, 2007, (72 FR 36482), Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742, made application by letter 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 |
|-----------------------|----------|
| Cocaine (9041) | II |
| Ecgonine (9180) | II |

The company plans on producing cocaine for sale to its customers, who are final dosage manufacturers. The ecgonine is formed during the manufacturing process for cocaine.

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 Johnson Matthey Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Johnson Matthey 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: November 6, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-22478 Filed 11-15-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 26, 2007, and published in the **Federal Register** on

July 5, 2007, (72 FR 36727), Abbott Laboratories, DBA Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey 07981, 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 I and II:

| Drug | Schedule |
|------------------------------|----------|
| Dihydromorphine (9145) | I |
| Hydromorphone (9150) | II |

The company plans to manufacture bulk product and dosage units 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 Abbott Laboratories to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Abbott Laboratories 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: November 5, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-22476 Filed 11-15-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 26, 2007, and published in the **Federal Register** on July 5, 2007, (72 FR 36729-36730), 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 schedule I and II: