Drug	Schedule
Fentanyl (9801)	II

The firm plans to manufacture small quantities of the listed controlled substances to produce isotope labeled standards for drug analysis.

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.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than November 3, 2003.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22328 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF justice

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Withdrawal of Application

As set forth in the **Federal Register** on April 2, 2003, (68 FR 16089), ISP Freetown Fine Chemicals, Inc., 238 South Main Street, Assonet, Massachusetts 02702, made application by letter to the Drug Enforcement Administration to be registered as a bulk manufacturer of Levorphanol (9220) a basic class of controlled substance listed in Schedule II.

By letter dated June 30, 2003, ISP Freetown Fine Chemicals, Inc., requested that their application to manufacture Levorphanol be withdrawn. Therefore, said application is hereby withdrawn.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22327 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 29, 2003 and published in the **Federal Register** on May 29, 2003, (68 FR 32088), Johnson Matthey, Inc., Pharmaceutical Materials, 2003 Nolte Drive West Deptford, New Jersey, 08066, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The firm plans to import Phenylacetone for conversion to amphetamine base to sell in bulk to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Johnson Matthey, Inc., to import the listed 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, at this time.

DEA has investigated Johnson Matthey, Inc., on a regular basis to ensure that the company's continued registration is consistent with the public interest. This investigation 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 Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 3101.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22326 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to section 1301.33(a) of title 21 of the Code of Federal Regulations (CFR), this is notice that on February 9, 2003, Lin-Zhi International, Inc., 687 North Pastoria Avenue, Sunnyvale, California 94085–2917, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxymethamphetamine (7405).	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Secrobarbital (2315)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Methadone (9250)	II
Dextropropoxyphene (9273)	II
Morphine (9300)	II

The firm plans to manufacture small quantities of controlled substances to make drug testing reagents and controls.

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.

Any such comments or objections may be addressed, in quituplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: Federal Register Representative, Officer of Chief Counsel (CCD) and must be filed no later than November 3, 2003.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22330 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated May 7, 2003, and published in the **Federal Register** on April 29, 2003, (68 FR 32088), Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Gamma hydroxybutyric acid (2010).	I

Drug	Schedule
Amphetamine (1100)	II
Phenylacetone (8501)	II
Methypenidate (1724)	II

The firm plans to produce bulk products for finished dosage units for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Lonza Riverside to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Lonza Riverside to ensure that the company's registration is consistent with the public interest. This 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed is granted.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22324 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances Notice of Registration

By Notice dated April 29, 2003, and published in the **Federal Register** on May 29, 2003, (68 FR 32089), Organichem Corporation, 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic class of Schedule II of controlled substance listed below:

Drug	Schedule
Dextropoxyphene (9273)	II

The firm plans to manufacture bulk products for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Organichem Corporation to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Organichem Corporation to ensure that the company's registration is consistent with the public interest. This 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed is granted.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22325 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated May 2, 2003 and published in the **Federal Register** on May 29, 2003, (68 FR 32089), Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Schedules I & II, for the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic Acid Diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Benzoylecogonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to import the listed controlled substances to manufacture diagnostic products for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Roche Diagnostics Corporation to import the listed 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, at this time. DEA has investigated Roche Diagnostics Corporation on a regular basis to ensure that the company's continued registration is consistent with the public interest. This investigation 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 Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22322 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated may 2, 2003, and published in the **Federal Register** on May 29, 2003, (68 FR 32089), Roche Diagnostics corporation, Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic Acid Diethylamide (7515)	I
Tetrahydrocannabinol (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Benzoylecogonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to manufacture small quantities of controlled substances for use in diagnostic products.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Rocke Diagnostics