371–6447. Written or faxed comments should be submitted by June 13, 2003.

Carol D. Shull,

Keeper of the National Register of Historic Places.

COLORADO

Douglas County

Devils Head Lookout, South Platte District, Pike National Forest, Sedalia, 03000518.

LOUISIANA

Orleans Parish

Broadmoor Historic District, Roughly bounded by South Broad/Fountainbleau, Milan, S. Claiborne and Octavia, New Orleans, 03000519.

MINNESOTA

Hennepin County

Lock and Dam No. 2, Mississippi R N of Lake St/Marshall Ave., Minneapolis, 03000522.

Le Sueur County

Dodd Road Discontiguous District, Roughly Cty Rd. 1 to MN 21, Cty Rd. 136 W of Kilkenny cont. NW to Cty Rd. 2, Cty Rd. 148 W of Cleveland., Forest, 03000520.

St. Louis County

Pyhala, Anna and Mikko, Farm, (Rural Finnish Log Buildings of St. Louis County, Minnesota, 1890–1930s MPS) 4745 Salo Rd., Embarrass, 03000521.

MISSOURI

Jackson County

Exchange Building, 1201–1207 Grand Blvd., Kansas City, 03000524.
Greenlease Cadillac Building, 2900 Gillham Rd., Kansas City, 03000523.
Knickerbocker Apartments, 501–535 Knickerbocker Place, Kansas City, 03000525.

NEW HAMPSHIRE

Grafton County

Greenleaf, Abbie, Library, 439 Main St., Franconia, 03000526.

PENNSYLVANIA

Delaware County

Booth Farm, 3221 Foulk Rd., Boothwyn, 03000527.

Philadelphia County

Philadelphia School of Occupational Therapy, 419 S. 19th St., Philadelphia, 03000528.

A request for REMOVAL has been made for the following resources:

MINNESOTA

Rice County

Church of St. Patrick—Catholic (Rice County MRA), Co. Hwy. 10 (Dodd Rd.), Faribault (vicinity), 82003032.

Scott County

Roehl-Lenzmeier House (Scott County MRA), MN 300 Shakopee (vicinity), 80002170.

St. Louis County

Hearding, John Harris, Grammar and High School and John A. Johnson Grammar School Jct. Of 4th Ave. N and First St. W, Aurora, 96001593.

[FR Doc. 03–13337 Filed 5–28–03; 8:45 am]

BILLING CODE 4312–51–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to section 1008 of the Controlled Substances Import and Export Act (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 section 1002(a) 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 section 1301.34 of title 21, Code of Federal Regulations (CFR), notice is hereby given that on June 25, 2002, Johnson Matthey, Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey, 08066, made application by renewal to the Drug Enforcement Administration 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.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing 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 (CCD), and must be filed no later than June 30, 2003.

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 at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import 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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: April 29, 2003.

Laura M. Nagel,

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

[FR Doc. 03–13312 Filed 5–28–03; 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 January 6, 2003, Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application by renewal and on April 14, 2003, by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Gamma hyrdoxybutyric acid (2010).	I
Amphetamine (1100)	II
Phenylacetone (8501)	II
Methyphenidate (1724)	II

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

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 July 28, 2003.

Dated: May 7, 2003.

Laura M. Nagel,

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

[FR Doc. 03–13309 Filed 5–28–03; 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 March 25, 2003, Organichem Corporation, 33 Riverside Avenue, Rensselaer, New York 12144, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basis 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.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objection 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 (CCD) and must be filed no later than July 28, 2002.

Dated: April 29, 2003.

Laura M. Nagel,

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

[FR Doc. 03–13311 Filed 5–28–03; 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 June 10, 2002, Organix, Inc., 240 Salem Street, Woburn, MA 01810, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic class of Cocaine (9041), a Schedule II controlled substance.

The firm plans to synthesize a controlled substance derivative from a non-controlled substance; the derivative will be sold to the firm's customer for research purposes.

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.

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: DEA Federal Register Representative (CCD) and must be filed no later than July 28, 2003.

Dated: May 7, 2003.

Laura M. Nagel,

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

[FR Doc. 03–13310 Filed 5–28–03; 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 January 28, 2003, Roche Diagnostics Corporation, Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal and on January 29, 2003, by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic Acid Diethylamide (7315)	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.

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 July 28, 2003.

Dated: May 2, 2003.

Laura M. Nagel,

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

[FR Doc. 03–13313 Filed 5–28–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances, Notice of Application

Pursuant to section 1008 of the Controlled Substances Import and Export Act (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 section 1002(a) 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 section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on January 28, 2003, Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration 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
Tetrahydrocannabinol (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.

Any manufacturer holding, or applying for, registration as a bulk