

releases or threatened releases of hazardous substances at the site, and the defendants will implement the remedy at the site.

The Department of Justice will receive comments relating to the proposed Partial Consent Decree for a period of 30 days from the date of this publication. Comments should be addressed to the Assistant Attorney General of the Environment and Natural Resources Division, Department of Justice, Washington, D.C. 20530. All comments should refer to *United States v.*

*Consolidated Rail Corp., et al.*, D.J. Ref. 90-11-3-594. Commenters may request an opportunity for a public meeting in the affected area, in accordance with Section 7003(d) of the Resource Conservation and Recovery Act, 42 U.S.C. § 6973(d).

The proposed Partial Consent Decree may be examined at the offices of the Environmental Protection Agency, 77 West Jackson Boulevard, Chicago, Illinois, 60604, or at the Consent Decree Library, 1120 G Street, N.W., 4th floor, Washington, D.C. 20005, 202-624-0892. A copy of the proposed Partial Consent Decree may be obtained in person or by mail from the Consent Decree Library. In requesting a copy (without attachments), please enclose a check in the amount of \$22.75 for the Decree (25 cents per page reproduction costs) payable to the Consent Decree Library. When requesting a copy, please refer to *United States v. Consolidated Rail Corp. et al.*, D.J. Ref. No. 90-11-3-594.

**Bruce S. Gelber,**

*Deputy Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

[FR Doc. 97-22599 Filed 8-25-97; 8:45am]

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## 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 regulation 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 May 29, 1997, Applied Science Labs, Inc., A Division of Altech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Heroin (9200) .....	I
Morphine (9300) .....	II

The firm plans to import these controlled substances for the manufacture of reference standards.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances 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: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

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 classes 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 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 21, 1997.

**John H. King,**

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

[FR Doc. 97-22559 Filed 8-25-97; 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 regulation 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 17, 1997, Bridgeway Trading Corporation, 7401 Metro Blvd., Suite 480, Minneapolis, Minnesota 55439, made application by renewal to the Drug Enforcement Administration to be registered as an importer of marihuana (7360) a basic class of controlled substance in Schedule I.

This application is exclusively for the importation of marihuana seed which will be rendered non-viable and used as bird seed.

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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

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 classes 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 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 22, 1997.

**John H. King,**

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

[FR Doc. 97-22560 Filed 8-25-97; 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 8, 1997, and published in the **Federal Register** on April 29, 1997, (62 FR 23268), Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of 4-Methoxyamphetamine (7411) a basic class of controlled substance listed Schedule I.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Celgene Corporation to manufacture 4-Methoxyamphetamine is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 C.F.R. §§ 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 above is granted.

Dated: July 29, 1997.

**John H. King,**

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

[FR Doc. 97-22561 Filed 8-25-97; 8:45 am]

BILLING CODE 4410-09-M

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated March 31, 1997, and published in the **Federal Register** on May 8, 1997 (62 FR 25209), Ganes Chemicals, Inc., Industrial Park Road, Pennsville, New Jersey 08070, 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 below:

Drug	Schedule
Methylphenidate (1724) .....	II
Amobarbital (2125) .....	II
Pentobarbital (2270) .....	II
Secobarbital (2315) .....	II
Glutethimide (2550) .....	II
Methadone (9250) .....	II
Methadone-intermediate (9254) ...	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Ganes Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 C.F.R. §§ 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 above is granted.

Dated: July 29, 1997.

**John H. King,**

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

[FR Doc. 97-22562 Filed 8-25-97; 8:45 am]

BILLING CODE 4410-09-M

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated February 26, 1997, and published in the **Federal Register** on March 19, 1997, (62 FR 13170), Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, 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 below:

Drug	Schedule
Tetrahydrocannabinols (7370) .....	I
Methylphenidate (1724) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Diprenorphine (9058) .....	II
Etorphine Hydrochloride (9059) ...	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II
Hydrocodone (9193) .....	II
Levorphanol (9220) .....	II

Drug	Schedule
Meperidine (9230) .....	II
Methadone (9250) .....	II
Methadone-intermediate (9254) ...	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Opium extracts (9610) .....	II
Opium fluid extract (9620) .....	II
Opium tincture (9630) .....	II
Opium powdered (9636) .....	II
Opium granulated (9640) .....	II
Levo-alphaacetylmethadol (9648) ..	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

DEA has considered the factors in Title 21, United States Code, Section 823(a), as well as information provided by other bulk manufacturers, and determined that the registration of Mallinckrodt Chemical, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 C.F.R. §§ 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 above is granted.

Dated July 28, 1997.

**John H. King,**

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

[FR Doc. 97-22563 Filed 8-25-97; 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 of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 10, 1997, Novartis Pharmaceutical Corp., 59 Route 10, East Hanover, New Jersey 07936, made application by letter dated July 10, 1997, to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the Schedule II controlled substance methylphenidate (1724).

The firm, which is currently registered with DEA as a bulk manufacturer of methylphenidate at another location plans to manufacture validation batches in preparation of