

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]

BILLING CODE 4410-15-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 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