

inspection during regular business hours, Monday through Friday within 30 days following the meeting.

Dated: January 4, 2001.

Lester A. Snow,

Regional Director, Mid-Pacific Region.

[FR Doc. 01-658 Filed 1-9-01; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated September 26, 2000, and published in the **Federal Register** on October 13, 2000, (65 FR 60976), B.I. Chemicals, Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, 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 the phenylacetone for the bulk manufacture of amphetamine.

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 B.I. Chemicals, Inc. to import phenylacetone 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 B.I. Chemicals, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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 class of controlled substance listed above.

Dated: December 11, 2000.

John H. King,

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

[FR Doc. 01-753 Filed 1-9-01; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application Correction

In the **Federal Register** (FR Doc. 00-30294) Vol. 65, No. 229 at page 70936, dated November 28, 2000, Cerilliant Corporation, 14050 Summit Drive, Suite 121, PO Box 80189, Austin, Texas 78708-0189, made application to the Drug Enforcement Administration to be registered as an importer for certain basic classes of controlled substances.

The listing of controlled substance should not have included marihuana (7350) or tilidine (9750). Therefore, the above listed controlled substances are hereby deleted from Cerilliant Corporation's application for registration as an importer.

The listing of controlled substance should have included etorphine (9056). Therefore, etorphine (9056) is hereby added to Cerilliant Corporation's application for registration as an importer.

The firm plans to import small quantities of the listed controlled substance for the manufacture of analytical reference standards.

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: 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 the 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 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: December 14, 2000.

John H. King,

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

[FR Doc. 01-758 Filed 1-9-01; 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 June 6, 2000, Chiragene, Inc., 7 Powder Horn Drive, Warren, New Jersey 07059, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
N-Ethylamphetamine (1475)	I
2,5-Dimethoxyamphetamine (7396).	I
3,4-Methylenedioxyamphetamine (7400).	I
4-Methoxyamphetamine (7411) ...	I
Amphetamine (1100)	II
Methylphenidate (1724)	II

The firm plans to manufacture the listed controlled substances to supply their 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: DEA Federal Register Representative (CCR), and must be filed no later than March 12, 2001.

Dated: December 4, 2000.

John H. King,

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

[FR Doc. 01-747 Filed 1-9-01; 8:45 am]

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