

Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Morphine (9300)	II

The firm plans to manufacture Tetrahydrocannabinols and a derivative of Morphine for use in diagnostic kits.

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 above application and may also file a written request for a hearing thereon in accordance with 21 CFR 1301.54 and in the form prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed 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 31, 1995.

Dated: February 17, 1995.

Gene R. Haislip,

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

[FR Doc. 95-4934 Filed 2-28-95; 8:45 am]

BILLING CODE 4410-09-M

Manufacturer of Controlled Substances; Registration

By Notice dated April 25, 1994, and published in the **Federal Register** on May 4, 1994, (59 FR 23082), Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Pholcodine (9314)	I
Alphacetylmethadol (9603)	I
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Benzoyllecgonine (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone-intermediate (9254) ...	II

Drug	Schedule
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 (9639)	II
Opium granulated (9640)	II
Oxymorphone (9652)	II
Phenazocine (9715)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

A written request for a hearing was not received, but a comment was filed by a registered manufacturer. The comment was considered, however, DEA determined that the application should be approved. Therefore, pursuant to section 303 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 and Title 21, Code of Federal Regulations, § 1301.54(e), 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: January 24, 1995.

Gene R. Haislip,

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

[FR Doc. 95-4937 Filed 2-28-95; 8:45 am]

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Importation of Controlled Substances; 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 § 1311.42 of title 21, Code of Federal Regulations (CFR), notice is hereby given that on December 13, 1994, Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application 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 is importing the Phenylacetone to manufacture Dextroamphetamine Sulfate.

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 1301.54 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed 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 1311.42 (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 a 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: February 17, 1995.

Gene R. Haislip,

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

[FR Doc. 95-4935 Filed 2-28-95; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-30,550]

Amended Certification Regarding Eligibility to Apply for Worker Adjustment Assistance

Grace Energy Corporation, Dallas, TX and Grace Petroleum Corporation Operating at the Following Locations: TA-W-30,550A Oklahoma, TA-W-30,550B Texas, TA-W-30,550C Alabama, TA-W-30,550D Colorado,