

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: January 17, 1995.

Gene R. Haislip,

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

[FR Doc. 95-1774 Filed 1-24-95; 8:45 am]

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Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 20, 1994, MD Pharmaceutical, Inc., 3501 West Garry Avenue, Santa Ana, California 92704, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methylphenidate (1724)	II
Diphenoxylate (9170)	II

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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 24, 1995.

Dated: January 17, 1995.

Gene R. Haislip,

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

[FR Doc. 95-1773 Filed 1-24-95; 8:45 am]

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Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 4, 1994, and published in the **Federal Register** on November 15, 1994, (59 FR 58857), Norac Company Inc., 405 S. Motor Avenue, Azusa, California 91702, made application to the Drug Enforcement Administration to be registered as a bulk manufacturer of Tetrahydrocannabinols (7370), a basic class of controlled substance listed in Schedule I.

No comments or objections have been received. 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), 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: January 17, 1995.

Gene R. Haislip,

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

[FR Doc. 95-1770 Filed 1-24-95; 8:45 am]

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Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 29, 1994, and published in the **Federal Register** on December 6, 1994, (59 FR 62750), Upjohn Company, 7171 Portage Road, M.L. 7011-126-5, Kalamazoo, Michigan 49001, made application to the Drug Enforcement Administration to be registered as a bulk manufacturer of 2,5-Dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

No comments or objections have been received. 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), 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: January 17, 1995.

Gene R. Haislip,

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

[FR Doc. 95-1771 Filed 1-24-95; 8:45 am]

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DEPARTMENT OF LABOR

Pension and Welfare Benefits Administration

Advisory Council on Employee Welfare and Pension Benefits Plan; Meeting

Pursuant to the authority contained in Section 512 of the Employee Retirement Income Security Act of 1974 (ERISA), 29 U.S.C. 1142, a public meeting of the Advisory Council on Employee Welfare and Pension Benefit Plans will be held on February 15, 1995, in Suite S-2508, U.S. Department of Labor Building, Third and Constitution Avenue NW., Washington, DC 20210.

The purpose of the meeting, which will begin at 9:30 a.m. is to consider the items listed below and to invite public comment on any aspect of the administration of ERISA.

- I. Welcome and Introduction of New Council Members
- II. Assistant Secretary's Report
 - A. PWBA Priorities for 1995
 - B. Report to Congress
 - C. Miscellaneous Issues
 - D. Announcement of Council Chairperson and Vice Chairperson
- III. Introduction of PWBA Senior Staff and Orientation of New Members
- IV. Report of Advisory Council Working Groups (1993/1994 Term)
- V. Determination of Council Working Group/s for 1995
- VI. Procedure for Establishing Council and Working Group Meeting Dates
- VII. Statements From the General Public
- VIII. Adjourn

Members of the public are encouraged to file a written statement pertaining to any topic concerning ERISA by submitting twenty (20) copies on or before February 10, 1995 to William E. Morrow, Executive Secretary, ERISA Advisory Council, U.S. Department Labor, Suite N-5677, 200 Constitution Avenue, NW., Washington, DC 20210. Individuals or representatives of organizations wishing to address the Advisory Council should forward their request to the Executive Secretary or telephone (202) 219-8753. Oral presentations will be limited to ten minutes, but an extended statement may be submitted for the record.

Organizations or individuals may also submit statements for the record without testifying. Twenty (20) copies of such statements should be sent to the Executive Secretary of the Advisory Council at the above address. Papers will be accepted and included in the record of the meeting if received on or before February 10, 1995.