The proposed consent decree may be examined at the United States Department of Justice, Environment and Natural Resources Division, 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, 1120 G Street, N.W., 4th Floor, Washington, D.C. 20005. In requesting a copy, please refer to the referenced case and enclose a check in the amount of $40.00 (25 cents per page reproduction costs), payable to the Consent Decree Library. Attachments to the proposed partial consent decree can be obtained for the additional amount of $31.00.

Joel M. Gross,

*Chief Environmental Enforcement Section.*

[FR Doc. 96–11221 Filed 5–3–96; 8:45 am]

**BILLING CODE 4410–01–M**

## Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated February 2, 1996, and published in the Federal Register on February 13, 1996, (61 FR 5571), Orpharm Inc., 728 West 19th Street, Houston, Texas 77008, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Methadone (9250) | II. |
| Methadone-intermediate (9254) | II. |
| Levo-alphacetylmethadol (9648) | II. |

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. § 823(a) and determined that the registration of Orpharm 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: April 29, 1996.

Gene R. Haislip,

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

[FR Doc. 96–11254 Filed 5–3–96; 8:45 am]

**BILLING CODE 4410–09–M**

### 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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on February 20, 1996, Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application, which was received for processing on April 10, 1996, to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Coca Leaves (9040) | II. |
| Opium, raw (9600) | II. |
| Opium poppy (9650) | II. |
| Opium Straw Concentrate (9670) | II. |

The firm plans to import the listed controlled substances for the manufacture of bulk pharmaceutical controlled substances.

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 1301.54 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 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 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 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: April 30, 1996.

Gene R. Haislip,

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

[FR Doc. 96–11215 Filed 5–3–96; 8:45 am]

**BILLING CODE 4410–09–M**

### 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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 27, 1996, Research Biochemicals, Limited Partnership, Attn: Richard Milius, 1–3 Strathmore Road, Natick, Massachusetts 01760, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Methaqualone (2565) | I |
| Ibogaine (7260) | I |
| Tetrahydrocannabinols (7370) | I |
| Bufotenine (7433) | I |
| Dimethyltryptamine (7435) | I |
| Etorphine (except HCl) (9056) | I |
| Methylphenidate (1724) | II |
| Pentobarbital (2270) | II |
| Diprenorphine (9058) | II |
| Etorphine Hydrochloride (9059) | II |
| Diphenoxylate (9170) | II |
| Metazocine (9240) | II |
| Methadone (9250) | II |
| Fentanyl (9801) | II |

The firm plans to import small quantities of the controlled substances to manufacture laboratory reference standards and neurochemicals.

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 1301.54 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 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 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 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: April 30, 1996.

Gene R. Haislip,

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

[FR Doc. 96–11217 Filed 5–3–96; 8:45 am]

**BILLING CODE 4410–09–M**

## 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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 14, 1996, Roberts Laboratories, Inc., 4 Industrial Way West, Eatontown, New Jersey 07724, made application to the Drug Enforcement Administration to be registered as an importer of propiram (9649) a basic class of controlled substance listed in Schedule I.

The firm plans to import the propiram to manufacture in bulk for product development.

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, 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 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: April 30, 1996.

Gene R. Haislip,

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

[FR Doc. 96–11216 Filed 5–3–96; 8:45 am]

**BILLING CODE 4410–09–M**

## Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 22, 1995, and published in the Federal Register on January 22, 1996, (61 FR 1604), Upjohn Company, 7171 Portage Road, M.L. 2000–41–109, 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. DEA has considered the factors in 21 U.S.C. 823(a) and determined that this registration of the Upjohn Company to manufacture the listed controlled substance is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistance Administration, 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: April 29, 1996.

Gene R. Haislip,

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

[FR Doc. 96–11255 Filed 5–3–96; 8:45 am]

**BILLING CODE 4410–09–M**

# DEPARTMENT OF LABOR

## Pension and Welfare Benefits Administration

**[Prohibited Transaction Exemption 96–30; Exemption Application No. D–09904, et al.]**

## Grant of Individual Exemptions; Aultman Retirement Savings Plan

**AGENCY:** Pension and Welfare Benefits Administration, Labor.

**ACTION:** Grant of individual exemptions.

**SUMMARY:** This document contains exemptions issued by the Department of Labor (the Department) from certain of the prohibited transaction restrictions of the Employee Retirement Income Security Act of 1974 (the Act) and/or the Internal Revenue Code of 1986 (the Code).

Notices were published in the Federal Register of the pendency before the Department of proposals to grant such exemptions. The notices set forth a summary of facts and representations contained in each application for exemption and referred interested persons to the respective applications for a complete statement of the facts and representations. The applications have been available for public inspection at the Department in Washington, D.C. The notices also invited interested persons to submit comments on the requested exemptions to the Department. In addition the notices stated that any interested person might submit a written request that a public hearing be held (where appropriate). The applicants have represented that they have complied with the requirements of the notification to interested persons. No public comments and no requests for a hearing, unless otherwise stated, were received by the Department.