

Dated: July 25, 1996.  
 Gene R. Haislip,  
*Deputy Assistant Administrator, Office of  
 Diversion Control, Drug Enforcement  
 Administration.*  
 [FR Doc. 96-19443 Filed 7-30-96; 8:45 am]  
**BILLING CODE 4410-09-M**

**Manufacturer of Controlled  
 Substances; Notice of Registration**

By Notice dated April 9, 1996, and published in the Federal Register on April 19, 1996, (61 FR 17322), Knoll Pharmaceuticals, 30 North Jefferson Road, Whippany, New Jersey 07981, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of hydromorphone (9150), a basic class of controlled substance listed in Schedule II.

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 Knoll Pharmaceuticals to manufacture hydromorphone 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 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: July 25, 1996.  
 Gene R. Haislip,  
*Deputy Assistant Administrator, Office of  
 Diversion Control, Drug Enforcement  
 Administration.*  
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**Importer of Controlled Substances;  
 Notice of Registration**

By Notice dated April 30, 1996, and published in the Federal Register on May 6, 1996, (61 FR 20275), Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application to the Drug Enforcement Administration (DEA) 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
Poppy Straw Concentrate (9670) .....	II

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 Penick Corporation to import the listed controlled substances 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. 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 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: July 25, 1996.  
 Gene R. Haislip,  
*Deputy Assistant Administrator, Office of  
 Diversion Control, Drug Enforcement  
 Administration.*  
 [FR Doc. 96-19445 Filed 7-30-96; 8:45 am]  
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**Importer of Controlled Substances;  
 Notice of Registration**

By Notice dated April 30, 1996, and published in the Federal Register on May 6, 1996, (61 FR 20276), Roberts Laboratories, Inc., 4 Industrial Way West, Eatontown, New Jersey 07724, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of propiram (9649), a basic class of controlled substance listed in Schedule I.

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 Roberts Laboratories, Inc. to import propiram 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. 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 1311.42, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: July 25, 1996.  
 Gene R. Haislip,  
*Deputy Assistant Administrator, Office of  
 Diversion Control, Drug Enforcement  
 Administration.*  
 [FR Doc. 96-19446 Filed 7-30-96; 8:45 am]  
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**Importer of Controlled Substances;  
 Notice of Registration**

By Notice dated May 22, 1996, and published in the Federal Register on May 30, 1996, (61 FR 27099), Roche Diagnostic Systems, Inc., 1080 U.S. Highway 202, Somerville, New Jersey 08876, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of tetrahydrocannabinols (7370), a basic class of controlled substance listed in Schedule I.

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 Roche Diagnostic Systems, Inc. to import tetrahydrocannabinols 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. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1311.42, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: July 25, 1996.  
 Gene R. Haislip,  
*Deputy Assistant Administrator, Office of  
 Diversion Control, Drug Enforcement  
 Administration.*  
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**Importer of Controlled Substances;  
 Notice of Registration**

By Notice dated May 22, 1996, and published in the Federal Register on May 30, 1996, (61 FR 27100), Wildlife Laboratories, Inc., 1401 Duff Drive, Suite 600, Ft. Collins, Colorado 80524, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Etorphine Hydrochloride (9059) ...	II
Carfentanil (9743) .....	II

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 Wildlife Laboratories, Inc. to import the listed controlled substances is consistent with the public interest and with United States