

(CFR), this is notice that on January 4, 2001, High Standard Products Corp., 14441 Beach Boulevard, #225, Westminster, California 92683, 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
Methaqualone (2565) .....	I
Lysergic acid diethylamide (7315) .....	I
Tetrahydrocannabinols (7370) ....	I
3,4-Methylenedioxymphetamine (7400) .....	I
3,4-Methylenedioxy-N-ethylamphetamine (7404) .....	I
3,4-Methylenedioxymphetamine (7405) .....	I
4-Methoxyamphetamine (7411) ..	I
Heroin (9200) .....	I
3-Methylfentanyl (9813) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Secobarbital (2315) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II
Hydrocodone (9193) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture analytical reference standards.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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 September 21, 2001.

Dated: July 13, 2001.

**Laura M. Nagel,**

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

[FR Doc. 01-18216 Filed 7-20-01; 8:45 am]

**BILLING CODE 4410-09-M**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 14, 2000, and published in the **Federal Register** on November 28, 2000 (65 FR 70937), ISP Freetown Chemicals, Inc., 238 South Main Street, Freetown, Massachusetts 02702, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396) .....	I
Amphetamine (1100) .....	II
Phenylacetone (8501) .....	II

The firm plans to block manufacture amphetamine for a customer and to bulk manufacture the phenylacetone for the manufacture of the amphetamine. the bulk 2,5-Dimethoxyamphetamine will be used for conversion into a Non-controlled substance.

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 ISP Freetown Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated ISP Freetown Chemicals, Inc. to ensure that the company's registration is consistent with the public interest.

This investigation included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and O.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 a basic classes of controlled substances listed above is granted.

Dated: July 13, 2001.

**Laura M. Nagel,**

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

[FR Doc. 01-18212 Filed 7-20-01; 8:45 am]

**BILLING CODE 4410-09-M**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 12, 2000, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, 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
Tetrahydrocannabinols (7370) .....	I
Difenoxin (9168) .....	I
Propriam (9649) .....	I
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II
Anileridine (9020) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Alfentanil (9737) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture the listed controlled substances in bulk to supply final dosage from manufacturers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comment 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 (60 days from publication).

Dated: July 13, 2001.

**Laura M. Nagel,**

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

[FR Doc. 01-18218 Filed 7-20-01; 8:45 am]

**BILLING CODE 4410-09-M**