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 May 13, 2002.

Dated: February 19, 2002.

Laura M. Nagel,

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

[FR Doc. 02–5792 Filed 3–11–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substance; Notice of Registration

By Notice dated July 13, 2001, and published in the **Federal Register** on July 23, 2001, (66 FR 38321), High Standard Products Corp., 14441 Beach Boulevard, #225, Westminster, California 92683, 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
Methaqualone (2565)	I
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxyamphetamine (7400)	I
3,4-Methylenedioxy-N-ethylamphetamine (7404)	I
3,4-Methylenedioxymethamphetamine (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.

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 High Standard Products Corp. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated High Standard Products Corp. to ensure that the company's registration is consistent with the public interest. This investigation has 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 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: February 19, 2002.

Laura M. Nagel,

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

[FR Doc. 02–5794 Filed 3–11–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 31, 2001, ISP Freetown Fine Chemicals, Inc., 238 South Main Street, Freetown, Massachusetts 02702, made application by renewal and by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Phenylacetone (8501)	II
Fentanyl (9801)	II

The firm plans to bulk manufacture amphetamine, methamphetamine, and fentanyl for customers 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.

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 May 13, 2002.

Dated: February 19, 2002.

Laura M. Nagel,

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

[FR Doc. 02–5793 Filed 3–11–02; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Market Street Market; Denial of Application

On or about August 27, 2001, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause (OTSC) by certified mail to Market Street Market (MSM), located in Chehalis, Washington, notifying it of an opportunity to show cause as to why the DEA should not deny its application, dated November 2, 1998, for a DEA Certificate of Registration as a distributor of the List I chemicals ephedrine, pseudoephedrine, and phenylpropanolamine, pursuant to 21 U.S.C. 823(h), as being inconsistent with the public interest. The order also notified MSM that, should no request for hearing be filed within 30 days, the right to a hearing would be waived.

The OTSC was received September 6, 2001, as indicated by the signed postal return receipt. Since that time, no further response has been received from the applicant nor any person purporting to represent the applicant. Therefore, the Administrator of the DEA, finding that (1) thirty days having passed since receipt of the Order to Show Cause, and (2) no request for a hearing having been received, concludes that MSM is deemed to have waived its right to a hearing. After considering relevant material from the investigative file in this matter, the Administrator now enters his final order without a hearing pursuant to 21 CFR 1301.43(d) and (e) and 1301.46.

The Administrator finds as follows. List I chemicals are chemicals that may be used in the manufacture of a controlled substance in violation of the