Attorney for the United States, U.S. Department of Justice, Antitrust Division, 325 Seventh Street, NW., Suite 300, Washington, DC 20530. (202) 353–4391.

[FR Doc. 03–3441 Filed 2–11–03; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances, Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 2, 2002, Cedarburg Pharmaceuticals, LLC, 870 Badger Circle, Grafton, Wisconsin 53024, 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
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II

The firm will manufacturer these controlled substances for distribution to its customers.

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 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 April 14, 2003.

Dated: February 5, 2003.

Laura M. Nagel,

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

[FR Doc. 03–3502 Filed 2–11–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated February 19, 2002, and published in the **Federal Register** on March 12, 2002 (67 FR 11142), ISP Freetown Fine Chemicals, Inc., 238 South Main Street, Freetown, Massachusetts 02702, made application by renewal and 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
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.

No comments or objections have been received. DEA has considered the factors in Title 21, U.S.C. section 823(a) and determined that the registration of ISP Freetown Fine 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 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 5, 2003.

Laura M. Nagel,

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

[FR Doc. 03–3503 Filed 2–11–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 23, 2002, and published in the **Federal Register** on September 5, 2002 (67 FR 58857), ISP Freetown Fine 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
Methylphenidate (1724)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II

The firm plans to bulk manufacture methylphenidate to produce a commercial product and manufacture the dextropropoxyphene to supply the generic market.

No comments or objections have been received. DEA has considered the factors in Title 21, U.S.C., section 823(a) and determined that the registration of ISP Freetown Fine 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 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 a basic classes of controlled substances listed above is granted.

Dated: February 5, 2003.

Laura M. Nagel,

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

[FR Doc. 03–3504 Filed 2–11–03; 8:45 am]

BILLING CODE 4410–09–M

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[03–012]

Notice of Information Collection

AGENCY: National Aeronautics and Space Administration (NASA).

ACTION: Notice of information collection.

SUMMARY: The National Aeronautics and Space Administration, as part of its continuing effort to reduce paperwork