

Drug	Schedule
Amobarbital (2125) .....	II
Pentobarbital (2270) .....	II
Secobarbital (2315) .....	II
Phenylacetone (8501) .....	II
Oxycodone (9143) .....	II

The firm plans to produce bulk product and finished dosage units 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: Drug Operations Section, Domestic Drug Unit (ODOD) and must be filed no later than June 2, 2003.

Dated: March 11, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-7825 Filed 4-1-03; 8:45 am]

**BILLING CODE 4410-09-M**

(ODOD) and must be filed no later than (June 25, 2003).

Dated: March 11, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-7830 Filed 4-1-03; 8:45 am]

**BILLING CODE 4410-09-M**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Registration

By Notice dated June 7, 2002, and published in the **Federal Register** on June 20, 2002, (67 FR 42059), Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

Objections and a request for hearing were timely filed and then withdrawn. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of ISP Freetown Fine Chemicals, Inc., 238 South Main Street, Asonet, Massachusetts 02702, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The firm plans to import the phenylacetone to manufacture amphetamine.

Objections and a request for hearing were timely filed and then withdrawn. DEA has considered the factors in Title 21, United States Code, 823(a) and determined that the registration of ISP Freetown Fine Chemicals, Inc., 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. DEA has investigated ISP Freetown Fine Chemicals, Inc. to ensure that the company's continued registration is consistent with the public interest. This investigation included inspection and testing of the company's physical security systems, certification of the company's compliance with state and local laws, and a review of the company's background and history. 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 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: March 14, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-7835 Filed 4-1-03; 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 17, 2002, ISP Freetown Fine Chemicals Inc., 238 South Main Street, Asonet, Massachusetts 02702, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Levorphanol (9220) a basic class of controlled substance listed in Schedule II.

The firm plans to produce bulk product providing an alternate supply of an active pharmaceutical ingredient to its customer.

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: Drug Operations Section, Domestic Drug Unit

Drug	Schedule
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Methylphenidate (1724) .....	II
Codeine (9050) .....	II
Codeine-N-oxide (9053) .....	I
Diprenorphine (9058) .....	II