

burden on the public is 1,082 hours annually.

If additional information is required contact: Brenda Dyer, Deputy Clearance Officer Information Management and Security Staff, Justice Management Division, United States Department of Justice, 601 D Street NW., Patrick Henry Building, Suite 1600, NW., Washington, DC 20530.

Dated: March 27, 2003.

Brenda Dyer,

Deputy Clearance Officer, Department of Justice.

[FR Doc. 03-7823 Filed 4-1-03; 8:45 am]

BILLING CODE 4410-AT-M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Import of Controlled Substances; Notice of Registration

By notice dated April 24, 2002, and published in the **Federal Register** on May 17, 2002 (67 FR 35136), Salsbury Chemicals, Inc., 1205 11th Street, Charles City, Iowa 50616-3466, 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's legal name has since changed to Cambrex Charles City, Inc.

The firm plans to import phenylacetone to manufacture amphetamine for distribution to its customers.

Objections and a request for hearing were timely filed and then withdrawn.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 Salsbury 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 Salsbury Chemicals, Inc. (now Cambrex Charles City, 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 system, verification 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.

Dated: March 14, 2003.

Laura M. Nagel,

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

[FR Doc. 03-7837 Filed 4-1-03; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated March 12, 2002, and published in the **Federal Register** on March 25, 2002 (67 FR 13664), Chiragene, Inc., 7 Powder Horn Drive, Warren, New Jersey 07059, made application by renewal 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 Company's legal name has since changed to Cambrex North Brunswick, Incorporated.

The firm plans to import 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, section 823(a) and determined that the registration of Cambrex North Brunswick, Inc., to import phenylacetone 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 Cambrex North Brunswick, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification 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.

Dated: March 14, 2003.

Laura M. Nagel,

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

[FR Doc. 03-7838 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 § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 22, 2003, Cedarburg Pharmaceuticals, LLC, 870 Badger Circle, Grafton, Wisconsin 53204, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of dihydromorphine (9145), a basic class of controlled substance listed in Schedule I.

The firm plans to use this substance in the conversion processes to produce Schedule II hydromorphone.

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 (ODOU) and must be filed no later than June 2, 2003.

Dated: March 14, 2003.

Laura M. Nagel,

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

[FR Doc. 03-7824 Filed 4-1-03; 8:45 am]

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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 22, 2002, October 9, 2002, and November 7, 2002, Cody Laboratories, Inc., 331 33rd Street, Cody, Wyoming 82414, made application by three separate letters to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic class of Schedule I and II controlled substances listed below:

Drug	Schedule
Dihydromorphine (9145)	I
Methamphetamine (1105)	II
Amphetamine (1100)	II

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]

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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, Assonet, 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

(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]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated February 19, 2002, and published in the **Federal Register** on March 5, 2002, (67 FR 9988), ISP Freetown Fine Chemicals, Inc., 238 South Main Street, Assonet, 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 Freeman 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]

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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 Lonza Riverside to import phenylacetone 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 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed.

Dated: March 14, 2003.

Laura M. Nagel,

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

[FR Doc. 03-7833 Filed 4-1-03; 8:45 am]

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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 January 14, 2003, Mallinckrodt, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substance listed below:

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