

Division, and either e-mailed to [pubcomment-ees.enrd@usdoj.gov](mailto:pubcomment-ees.enrd@usdoj.gov) or mailed to P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044-7611, and should refer to *United States of America v. El Dorado County, California, et al*, Civil No. S-01-1520 MCE GGH (E.D. Cal.) (DOJ Ref. No. 90-11-3-06554) (Partial Consent Decree with STR).

The Partial Consent Decree with STR may be examined at U.S. Department of Agriculture, Office of General Counsel, 33 New Montgomery Street, 17th Floor, San Francisco, CA 94150 (contact Rose Mikovsky, (415) 744-3158). During the public comment period, the Partial Consent Decree with STR may also be examined on the following Department of Justice Web site, [http://www.usdoj.gov/enrd/Consent\\_Decrees.html](http://www.usdoj.gov/enrd/Consent_Decrees.html). A copy of the Partial Consent Decree with STR may also be obtained by mail from the Consent Decree Library, U.S. Department of Justice, P.O. Box 7611, Washington, DC 20044-7611 or by faxing or e-mailing a request to Tonia Fleetwood ([tonia.fleetwood@usdoj.gov](mailto:tonia.fleetwood@usdoj.gov)), fax no. (202) 514-0097, phone confirmation number (202) 514-1547. In requesting a copy from the Consent Decree Library, please refer to *United States of America v. El Dorado County, California, et al*, Civil No. S-01-1520 MCE GGH (E.D. Cal.) (DOJ Ref. No. 90-11-3-06554) (Partial Consent Decree with STR), and enclose a check in the amount of \$6.00 (25 cents per page reproduction cost) payable to the U.S. Treasury or, if by email or fax, forward a check in that amount to the Consent Decree Library at the stated address.

**Maureen Katz,**

*Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

[FR Doc. 2010-4310 Filed 3-2-10; 8:45 am]

**BILLING CODE 4410-15-P**

**DEPARTMENT OF JUSTICE**

**Office of Justice Programs**

[OJP (NIJ) Docket No. 1512]

**Draft NIJ Restraints Standard for Criminal Justice**

**AGENCY:** National Institute of Justice, Office of Justice Programs, DOJ.

**ACTION:** Notice of Draft NIJ Restraints Standard for Criminal Justice and Certification Program Requirements.

**SUMMARY:** In an effort to obtain comments from interested parties, the U.S. Department of Justice, Office of Justice Programs, National Institute of

Justice will make available to the general public two draft documents: (1) A draft standard entitled, "NIJ Restraints Standard for Criminal Justice" and (2) a draft companion document entitled, "NIJ Restraints Certification Program Requirements". The opportunity to provide comments on these two documents is open to industry technical representatives, criminal justice agencies and organizations, research, development and scientific communities, and all other stakeholders and interested parties. Those individuals wishing to obtain and provide comments on the draft documents under consideration are directed to the following Web site: <http://www.justnet.org>.

**DATES:** Comments must be received on or before April 19, 2010.

**FOR FURTHER INFORMATION CONTACT:** Casandra Robinson, by telephone at 202-305-2596 [Note: this is not a tollfree telephone number], or by e-mail at [casandra.robinson@usdoj.gov](mailto:casandra.robinson@usdoj.gov).

**Kristina Rose,**

*Acting Director, National Institute of Justice.*

[FR Doc. 2010-4434 Filed 3-2-10; 8:45 am]

**BILLING CODE 4410-18-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration**

By Notice dated November 23, 2009, and published in the **Federal Register** on December 2, 2009 (74 FR 63156), ISP Freetown Fine Chemicals, 238 South Main Street, Assonet, Massachusetts 02702, 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 plans to import the controlled substance to manufacture amphetamine.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of ISP Freetown Fine Chemicals to import the basic class of controlled substance 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 to ensure that the company's registration is consistent with the public interest. The 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: February 25, 2009.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2010-4399 Filed 3-2-10; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration**

By Notice dated November 20, 2009, and published in the **Federal Register** on November 30, 2009 (74 FR 62598), Aptuit, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Poppy Straw Concentrate (9670), a basic class of controlled substance listed in schedule II.

The company plans to import an ointment for the treatment of wounds which contain trace amounts of the controlled substance normally found in poppy straw concentrate for packaging and labeling for clinical trials.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Aptuit to import the basic class of controlled substance 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 Aptuit to ensure that the company's registration is consistent with the public interest. The 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: February 24, 2010.

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

[FR Doc. 2010-4440 Filed 3-2-10; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances;  
Notice of Registration**

By Notice dated October 16, 2009, and published in the **Federal Register** on October 28, 2009 (74 FR 55584), Clinical Supplies Management, 342 42nd Street South, Fargo, North Dakota 58103, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Poppy Straw Concentrate (9670), a basic class of controlled substance listed in schedule II.

The company plans to import an ointment for the treatment of wounds which contains trace amounts of controlled substances normally found in poppy straw concentrate which will be packaged and labeled for clinical trials.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and § 952(a), and determined that the registration of Clinical Supplies Management to import the basic class of controlled substance 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 Clinical Supplies Management to ensure that the company's registration is consistent with the public interest. The 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. 952(a) and § 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: February 24, 2010.

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

[FR Doc. 2010-4439 Filed 3-2-10; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled  
Substances; Notice of Registration**

By Notice dated October 30, 2009, and published in the **Federal Register** on November 6, 2009, (74 FR 57522), ISP Freetown Fine Chemicals, 238 South Main Street, Assonet, Massachusetts 02702, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396) .....	I
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II
Phenylacetone (8501) .....	II
Dextropropoxyphene, bulk (non-dosage forms) (9273) .....	II

The company plans to manufacture bulk API, for distribution to its customers. The bulk 2,5-Dimethoxyamphetamine will be used for conversion into non-controlled substances.

No comments or objections have been received. DEA has considered the factors in 21 USC 823(a) and determined that the registration of ISP Freetown Fine Chemicals to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated ISP Freetown Fine Chemicals to ensure that the company's registration is consistent with the public interest. The 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 USC 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 25, 2010.

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

[FR Doc. 2010-4400 Filed 3-2-10; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled  
Substances; Notice of Registration**

By Notice dated October 16, 2009, and published in the **Federal Register** on October 28, 2009, (74 FR 55587), Cayman Chemical Company, 1180 East Ellsworth Road, Ann Arbor, Michigan 48108, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedule I:

Drug	Schedule
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	I

The company plans to manufacture small quantities of marihuana derivatives for research purposes.

In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol. In reference to drug code 7370 (Tetrahydrocannabinols), the company will manufacture a synthetic THC. No other activity for this drug code is authorized for this registration.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cayman Chemical Company to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cayman Chemical Company to ensure that the company's registration is consistent with the public interest. The 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 24, 2010.

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

[FR Doc. 2010-4401 Filed 3-2-10; 8:45 am]

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