

study and install process changes or controls to eliminate excess emissions, and comply with interim emission limits.

The Department of Justice will receive comments relating to the proposed Consent Decree for a period of thirty (30) days from the date of this publication. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources 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 v. JELD-WEN, Inc.*, DJ Ref. No. 90-5-2-1-09567.

The proposed Consent Decree may be examined at the Office of the United States Attorney for the District of Oregon, 1000 SW. Third Avenue, Suite 600, Portland, Oregon 97204-2902, 503-727-1053, and at the Environmental Protection Agency, Region 10, 1200 6th Avenue, Seattle, Washington 98101,

800-424-4372. During the public comment period, the proposed Agreement may also be examined on the following Department of Justice Web site, <http://www.usdoj.gov/enrd/Consent-Decrees.html>. A copy of the proposed Agreement may also be obtained by mail from the Consent Decree Library, P.O. Box 7611, U.S. Department of Justice, 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 enclose a check in the amount of \$18.00 (25 cents per page reproduction cost) payable to the U.S. Treasury.

**Robert E. Maher, Jr.,**

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

[FR Doc. 2011-10052 Filed 4-26-11; 8:45 am]

**BILLING CODE 4410-15-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

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

By Notice dated February 15, 2011, and published in the **Federal Register** on February 23, 2011, 76 FR 10067, Sigma Aldrich Manufacturing LLC., 3500 Dekalb Street, St. Louis, Missouri 63118, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Cathinone (1235) .....	I
Methcathinone (1237) .....	I
Aminorex (1585) .....	I
Gamma Hydroxybutyric Acid (2010) .....	I
Methaqualone (2565) .....	I
Alpha-ethyltryptamine (7249) .....	I
Ibogaine (7260) .....	I
Lysergic acid diethylamide (7315) .....	I
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	I
Mescaline (7381) .....	I
4-Bromo-2,5-dimethoxyamphetamine (7391) .....	I
4-Bromo-2,5-dimethoxyphenethylamine (7392) .....	I
4-Methyl-2,5-dimethoxyamphetamine (7395) .....	I
2,5-Dimethoxyamphetamine (7396) .....	I
3,4-Methylenedioxyamphetamine (7400) .....	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402) .....	I
3,4-Methylenedioxy-N-ethylamphetamine (7404) .....	I
3,4-Methylenedioxymethamphetamine (MDMA) (7405) .....	I
4-Methoxyamphetamine (7411) .....	I
Bufotenine (7433) .....	I
Diethyltryptamine (7434) .....	I
Dimethyltryptamine (7435) .....	I
Psilocybin (7437) .....	I
Psilocyn (7438) .....	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470) .....	I
N-Benzylpiperazine (BZP) (7493) .....	I
Heroin (9200) .....	I
Normorphine (9313) .....	I
Etonitazene (9624) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Methylphenidate (1724) .....	II
Amobarbital (2125) .....	II
Pentobarbital (2270) .....	II
Secobarbital (2315) .....	II
Glutethimide (2550) .....	II
Nabilone (7379) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Diprenorphine (9058) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II

Drug	Schedule
Ecgonine (9180) .....	II
Ethylmorphine (9190) .....	II
Hydrocodone (9193) .....	II
Levorphanol (9220) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Opium, powdered (9639) .....	II
Levo-alphaacetylmethadol (9648) .....	II
Oxymorphone (9652) .....	II
Fentanyl (9801) .....	II

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis.

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 Sigma Aldrich Manufacturing LLC. to import the basic classes of controlled substances 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 Sigma Aldrich Manufacturing LLC. 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 classes of controlled substances listed.

Dated: April 15, 2011.

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

[FR Doc. 2011-10145 Filed 4-26-11; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Application**

Pursuant to § 1301.33(a) Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 31, 2011, Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, made application by letter to the Drug Enforcement Administration

(DEA) to be registered as a bulk manufacturer of 4-Anilino-N-phenethyl-4-Piperidine (8333), a basic class of controlled substance listed in schedule II.

The company plans to use this controlled substance in the manufacture of another controlled substance.

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 pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than June 27, 2011.

Dated: April 15, 2011.

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

[FR Doc. 2011-10139 Filed 4-26-11; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

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

By Notice dated October 6, 2010, and published in the **Federal Register** on October 14, 2010, 75 FR 63203, PCAS-Nanosyn, LLC, 3331-B Industrial Drive, Santa Rosa, California 95403, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Methylphenidate (1724) .....	II
Phencyclidine (7471) .....	II
Codeine (9050) .....	II

Drug	Schedule
Diprenorphine (9058) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Oxymorphone (9652) .....	II
Fentanyl (9801) .....	II

The company is a contract manufacturer. At the request of the company's customers, it manufactures derivatives of controlled substances in bulk form only. The primary service provided by the company to its customers is the development of the process of manufacturing the derivative. As part of its service to its customers, the company distributes the derivatives of the controlled substances it manufactures to those customers. The company's customers use the newly-created processes and the manufactured derivatives in furtherance of formulation processes and dosage form manufacturing; pre-clinical studies, including toxicological studies; clinical studies supporting investigational Drug Applications; and use in stability studies.

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 PCAS-Nanosyn, LLC to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated PCAS-Nanosyn, LLC 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of