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.enu@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., D 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 Website, 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), 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.
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 another controlled substance.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

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).

The company plans to ensure that 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 952(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. 952(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of