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-alphacetylmethadol (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
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

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. 2011–10144 Filed 4–26–11; 8:45 am]

BILLING CODE 4410–09–P

NUCLEAR REGULATORY COMMISSION

[Docket No. NRC–2010–0377]

Agency Information Collection Activities: Submission for the Office of Management and Budget (OMB) Review; Comment Request

AGENCY: Nuclear Regulatory Commission (NRC).

ACTION: Notice of the OMB review of information collection and solicitation of public comment.

SUMMARY: The NRC has recently submitted to OMB for review the following proposal for the collection of information under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). The NRC hereby informs potential respondents that an agency may not conduct or sponsor, and that a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The NRC published a **Federal Register** Notice with a 60-day comment period on this information collection on December 23, 2010.

1. *Type of submission, new, revision, or extension:* Extension.

2. *The title of the information collection:* NUREG/BR–0238, Materials Annual Fee Billing Handbook; NRC Form 628, "Financial EDI Authorization;" NUREG/BR–0254, Payment Methods; and NRC Form 629, "Authorization for Payment by Credit Card."

3. *Current OMB approval number:* 3150–0190.

4. *The form number if applicable:* NRC Form 628, "Financial EDI Authorization" and NRC Form 629, "Authorization for Payment by Credit Card.

5. *How often the collection is required:* On occasion (as needed to pay invoices).

6. *Who will be required or asked to report:* Anyone doing business with the Nuclear Regulatory Commission including licensees, applicants and individuals who are required to pay a fee for inspections and licenses.

7. *An estimate of the number of annual responses:* 583 (11 for NRC form 628 and 572 for NRC form 629 and NUREG/BR–0254).

8. *The estimated number of annual respondents:* 583 (11 for NRC form 628 and 572 for NRC form 629 and NUREG/BR–0254).

9. *An estimate of the total number of hours needed annually to complete the requirement or request:* 47 hours (.9 hour for NRC form 628 and 46 hours for NRC form 629 and NUREG/BR–0254).

10. *Abstract:* The U.S. Department of the Treasury encourages the public to pay monies owed the government through use of the Automated Clearinghouse Network and credit cards. These two methods of payment are used by licensees, applicants, and individuals to pay civil penalties, full cost licensing fees, and inspection fees to the NRC.

The public may examine and have copied for a fee publicly available documents, including the final supporting statement, at the NRC's Public Document Room, Room O–1F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852. OMB clearance requests are available at the NRC worldwide Web site: *http://www.nrc.gov/public-involve/doc-comment/omb/*. The document will be available on the NRC home page site for 60 days after the signature date of this notice.

Comments and questions should be directed to the OMB reviewer listed below by May 27, 2011. Comments received after this date will be considered if it is practical to do so, but assurance of consideration cannot be given to comments received after this date.

Christine J. Kymn, Desk Officer, Office of Information and Regulatory Affairs (3150–0190), NEOB–10202, Office of Management and Budget, Washington, DC 20503.

Comments can also be e-mailed to *Christine_J._Kymn@omb.eop.gov* or submitted by telephone at 202–395–4638.

The NRC Clearance Officer is Tremaine Donnell, 301–415–6258.

Dated at Rockville, Maryland, this 21st day of April, 2011.

For the Nuclear Regulatory Commission.

Tremaine Donnell,

NRC Clearance Officer, Office of Information Services.

[FR Doc. 2011–10162 Filed 4–26–11; 8:45 am]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

[Docket No. NRC–2011–0056]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Nuclear Regulatory Commission (NRC).

ACTION: Notice of pending NRC action to submit an information collection request to the Office of Management and Budget (OMB) and solicitation of public comment.

SUMMARY: The NRC invites public comment about our intention to request the OMB's approval for renewal of an existing information collection that is summarized below. We are required to publish this notice in the **Federal Register** under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

Information pertaining to the requirement to be submitted:

1. *The title of the information collection:* 10 CFR part 81, "Standard Specifications for Granting of Patent Licenses."

2. *Current OMB approval number:* 3150–0121.

3. *How often the collection is required:* Applications for licenses are submitted once. Other reports are submitted annually or as other events require.

4. *Who is required or asked to report:* Applicants for and holders of NRC licenses to NRC inventions.

5. *The number of annual respondents:* 1.

6. *The number of hours needed annually to complete the requirement or request:* 37; however, no applications are anticipated during the next 3 years.

7. *Abstract:* As specified in 10 CFR part 81, the NRC may grant non-exclusive licenses or limited exclusive licenses to its patent inventions to responsible applicants. Applicants for licenses to NRC inventions are required to provide information which may provide the basis for granting the requested license. In addition, all license holders must submit periodic reports on efforts to bring the invention to a point of practical application and the extent to which they are making the benefits of the invention reasonably accessible to the public. Exclusive license holders must submit additional information if they seek to extend their licenses, issue sublicenses, or transfer the licenses. In addition, if requested, exclusive license holders must promptly supply to the United States Government copies of all pleadings and other papers