

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]

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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 Morrisette 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

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/](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