

the basic classes of controlled substances listed.

Dated: January 16, 2014.

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

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration; Clinical Supplies Management, Inc.**

By Notice dated August 29, 2013, and published in the **Federal Register** on September 6, 2013, 78 FR 54913, Clinical Supplies Management, Inc., 342 42nd Street South, Fargo, North Dakota 58103, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Sufentanil (9740), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance with the sole purpose of packaging, labeling, and distributing to customers which are qualified clinical sites, conducting FDA-approved 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, Inc., 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. DEA has investigated Clinical Supplies Management, Inc., 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: January 14, 2014.

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

[FR Doc. 2014-01788 Filed 1-29-14; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration; Halo Pharmaceutical, Inc.**

By Notice dated August 14, 2013, and published in the **Federal Register** on August 20, 2013, 78 FR 51210, Halo Pharmaceutical, Inc., 30 North Jefferson Road, Whippany, New Jersey 07981, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Dihydromorphine (9145) .....	I
Hydromorphone (9150) .....	II

Dihydromorphine is an intermediate in the manufacture of Hydromorphone and is not for commercial distribution.

The company plans to manufacture Hydromorphone for sale to other manufacturers and to manufacture other controlled substances for distribution to its customers.

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 Halo Pharmaceutical, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Halo Pharmaceutical, Inc., 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: January 15, 2014.

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

[FR Doc. 2014-01785 Filed 1-29-14; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration; Chattem Chemicals, Inc.**

By Notice dated August 14, 2013, and published in the **Federal Register** on August 20, 2013, 78 FR 51210, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
4-Methoxyamphetamine (7411) ...	I
Dihydromorphine (9145) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Lisdexamfetamine (1205) .....	II
Methylphenidate (1724) .....	II
Pentobarbital (2270) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Methadone intermediate (9254) ...	II
Morphine (9300) .....	II
Oripavine (9330) .....	II
Thebaine (9333) .....	II
Opium tincture (9630) .....	II
Opium, powdered (9639) .....	II
Opium, granulated (9640) .....	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Remifentanil (9739) .....	II
Sufentanil (9740) .....	II
Tapentadol (9780) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers. Regarding (9640), the company plans to manufacture another controlled substance for sale to its customers.

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 Chattem Chemicals, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Chattem Chemicals, Inc., 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: January 15, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014-01782 Filed 1-29-14; 8:45 am]

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## DEPARTMENT OF LABOR

### Office of the Secretary

#### **Agency Information Collection Activities; Submission for OMB Review; Comment Request; Application for Approval of a Representative's Fee in Black Lung Claim Proceedings Conducted by the U.S. Department of Labor**

**ACTION:** Notice.

**SUMMARY:** The Department of Labor (DOL) is submitting the Office of Workers' Compensation Programs sponsored information collection request (ICR) revision titled, "Application for Approval of a Representative's Fee in Black Lung Claim Proceedings Conducted by the U.S. Department of Labor," to the Office of Management and Budget (OMB) for review and approval for use in accordance with the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501 et seq.).

**DATES:** Submit comments on or before March 3, 2014.

**ADDRESSES:** A copy of this ICR with applicable supporting documentation; including a description of the likely respondents, proposed frequency of response, and estimated total burden may be obtained free of charge from the RegInfo.gov Web site at [http://www.reginfo.gov/public/do/PRAViewICR?ref\\_nbr=201309-1240-002](http://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=201309-1240-002) (this link will only become active on the day following publication of this notice) or by contacting Michel Smyth by telephone at 202-693-4129, TTY 202-693-8064, (these are not toll-free numbers) or sending an email to [DOL\\_PRA\\_PUBLIC@dol.gov](mailto:DOL_PRA_PUBLIC@dol.gov).

Submit comments about this request by mail or courier to the Office of Information and Regulatory Affairs,

Attn: OMB Desk Officer for DOL-OWCP, Office of Management and Budget, Room 10235, 725 17th Street NW., Washington, DC 20503; by Fax: 202-395-6881 (this is not a toll-free number); or by email: [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov). Commenters are encouraged, but not required, to send a courtesy copy of any comments by mail or courier to the U.S. Department of Labor—OASAM, Office of the Chief Information Officer, Attn: Departmental Information Compliance Management Program, Room N1301, 200 Constitution Avenue NW., Washington, DC 20210; or by email: [DOL\\_PRA\\_PUBLIC@dol.gov](mailto:DOL_PRA_PUBLIC@dol.gov).

**FOR FURTHER INFORMATION CONTACT:**

Michel Smyth by telephone at 202-693-4129, TTY 202-693-8064, (these are not toll-free numbers) or sending an email to [DOL\\_PRA\\_PUBLIC@dol.gov](mailto:DOL_PRA_PUBLIC@dol.gov).

**Authority:** 44 U.S.C. 3507(a)(1)(D).

**SUPPLEMENTARY INFORMATION:** This ICR seeks OMB approval under the PRA to revise the Application for Approval of a Representative's Fee in Black Lung Claim Proceedings Conducted by the U.S. Department of Labor, Form CM-972, to include information about how a respondent with a disability may contact the OWCP in order to receive assistance in completing the form. There is no change to the information collected from the public.

Individuals filing with the OWCP, Division of Coal Mine Workers' Compensation for benefits under the Black Lung Benefits Act may elect to be represented or assisted by an attorney or other representative. For those cases that are approved, 30 U.S.C. 901 of the Black Lung Benefits Act and 20 CFR 725.365-6 of the Black Lung Regulations have established standards for the information and documentation that must be submitted to the program for review so that the representative may be paid for services rendered to the claimant. Upon receipt of that evidence, the adjudicating official is required by regulation to evaluate the application and, based on the supporting information in the claim file, approve a fee for services rendered. Form CM-972 provides a standardized format that assists representatives participating in the Black Lung Benefits Program to submit the required information.

Form CM-972 is sent to and completed by the authorized representative of a black lung claimant whose claim has been approved for benefits. The completed form is then returned to and evaluated by the district director, administrative law judge, or appropriate appellate tribunal before whom the claimed services were

performed, and a fee amount is determined.

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless it is approved by the OMB under the PRA and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid Control Number. See 5 CFR 1320.5(a) and 1320.6. The DOL obtains OMB approval for this information collection under Control Number 1240-0011. The current approval is scheduled to expire on January 31, 2014; however, the DOL notes that existing information collection requirements submitted to the OMB receive a month-to-month extension while they undergo review. New requirements would only take effect upon OMB approval. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on September 11, 2013 (77 FR 55761).

Interested parties are encouraged to send comments to the OMB, Office of Information and Regulatory Affairs at the address shown in the **ADDRESSES** section within 30 days of publication of this notice in the **Federal Register**. In order to help ensure appropriate consideration, comments should mention OMB Control Number 1240-0011. The OMB is particularly interested in comments that:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

*Agency:* DOL-OWCP.

*Title of Collection:* Application for Approval of a Representative's Fee in