

the following basic classes of controlled substances:

Drug	Schedule
Tetrahydrocannabinols (7370) .....	I
Ecgonine (9180) .....	II
Morphine (9300) .....	II

The company plans to produce the listed controlled substances in bulk to be used in the manufacture of reagents and drug calibrator/controls which are DEA exempt products.

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 July 5, 2011.

Dated: April 25, 2011.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2011-10861 Filed 5-3-11; 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 January 5, 2011, Johnson Matthey Pharma Services, 70 Flagship Drive, North Andover, Massachusetts 01845, 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
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II
Hydrocodone (9193) .....	II

The company plans to utilize this facility to manufacture small quantities of the listed controlled substances in bulk and to conduct analytical testing in support of the company's primary manufacturing facility in West Deptford, New Jersey. The controlled substances manufactured in bulk at this facility will be distributed to the company's customers.

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 July 5, 2011.

Dated: April 28, 2011.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2011-10913 Filed 5-3-11; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

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

By Notice dated January 26, 2011, and published in the **Federal Register** on February 3, 2011, 76 FR 6159, Johnson Matthey Inc., Pharmaceuticals Materials, 900 River Road, Conshohocken, Pennsylvania 19428, made application by letter 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
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers. The Thebaine (9333) will also be used to manufacture other controlled substances for sale in bulk 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 Johnson Matthey Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Johnson Matthey 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: April 26, 2011.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2011-10865 Filed 5-3-11; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF LABOR**

**Occupational Safety and Health Administration**

[Docket No. OSHA-2011-0059]

**Occupational Exposure to Hazardous Chemicals in Laboratories Standard; Extension of the Office of Management and Budget's (OMB) Approval of Information Collection (Paperwork) Requirements**

**AGENCY:** Occupational Safety and Health Administration (OSHA), Labor.

**ACTION:** Request for public comments.

**SUMMARY:** OSHA solicits public comments concerning its proposal to extend OMB approval of the

information collection requirements specified in the Standard on Occupational Exposure to Hazardous Chemicals in Laboratories.

**DATES:** Comments must be submitted (postmarked, sent, or received) by July 5, 2011.

**ADDRESSES:**

*Electronically:* You may submit comments and attachments electronically at <http://www.regulations.gov>, which is the Federal eRulemaking Portal. Follow the instructions online for submitting comments.

*Facsimile:* If your comments, including attachments, are not longer than 10 pages, you may fax them to the OSHA Docket Office at (202) 693-1648.

*Mail, hand delivery, express mail, messenger, or courier service:* When using this method, you must submit your comments and attachments to the OSHA Docket Office, Docket No. OSHA-2011-0059, U.S. Department of Labor, Room N-2625, 200 Constitution Avenue, NW., Washington, DC 20210. Deliveries (hand, express mail, messenger, and courier service) are accepted during the Department of Labor's and Docket Office's normal business hours, 8:15 a.m. to 4:45 p.m., e.t.

*Instructions:* All submissions must include the Agency name and OSHA docket number (OSHA-2011-0059) for the Information Collection Request (ICR). All comments, including any personal information you provide, are placed in the public docket without change, and may be made available online at <http://www.regulations.gov>. For further information on submitting comments see the "Public Participation" heading in the section of this notice titled **SUPPLEMENTARY INFORMATION**.

*Docket:* To read or download comments or other material in the docket, go to <http://www.regulations.gov> or the OSHA Docket Office at the address above. All documents in the docket (including this **Federal Register** notice) are listed in the <http://www.regulations.gov> index; however, some information (e.g., copyrighted material) is not publicly available to read or download through the Web site. All submissions, including copyrighted material, are available for inspection and copying at the OSHA Docket Office. You also may contact Todd Owen at the address below to obtain a copy of the ICR.

**FOR FURTHER INFORMATION CONTACT:** Todd Owen, Directorate of Standards and Guidance, OSHA, U.S. Department of Labor, Room N-3609, 200

Constitution Avenue, NW., Washington, DC 20210; telephone (202) 693-2222.

**SUPPLEMENTARY INFORMATION:**

**1. Background**

The Department of Labor, as part of its continuing effort to reduce paperwork and respondent (i.e., employer) burden, conducts a preclearance consultation program to provide the public with an opportunity to comment on proposed and continuing information collection requirements in accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. 3506(c)(2)(A)).

This program ensures that information is in the desired format, reporting burden (time and costs) is minimal, collection instruments are clearly understood, and OSHA's estimate of the information collection burden is accurate. The Occupational Safety and Health Act of 1970 (the OSH Act) (29 U.S.C. 651 *et seq.*) authorizes information collection by employers as necessary or appropriate for enforcement of the OSH Act or for developing information regarding the causes and prevention of occupational injuries, illnesses, and accidents (29 U.S.C. 657). The OSH Act also requires that OSHA obtain such information with minimum burden upon employers, especially those operating small businesses, and to reduce to the maximum extent feasible unnecessary duplication of efforts in obtaining information (29 U.S.C. 657).

The Standard entitled "Occupational Exposure to Hazardous Chemicals in Laboratories" (29 CFR 1910.1450; the "Standard") applies to laboratories that use hazardous chemicals in accordance with the Standard's definitions for "laboratory use of hazardous chemicals" and "laboratory scale." The Standard requires these laboratories to maintain worker exposures at or below the permissible exposure limits specified for the hazardous chemicals in 29 CFR part 1910, subpart Z. They do so by developing a written Chemical Hygiene Plan (CHP) that describes: Standard operating procedures for using hazardous chemicals; hazard-control techniques; equipment-reliability measures; worker information-and-training programs; conditions under which the employer must approve operations, procedures, and activities before implementation; and medical consultations and examinations. The CHP also designates personnel responsible for implementing the CHP, and specifies the procedures used to provide additional protection to workers exposed to particularly hazardous chemicals.

Other information collection requirements of the Standard include: Documenting exposure monitoring results; notifying workers in writing of these results; presenting specified information and training to workers; establishing a medical surveillance program for overexposed workers; providing required information to the physician; obtaining the physician's written opinion on using proper respiratory equipment; and, establishing, maintaining, transferring, and disclosing exposure monitoring and medical records. These collection of information requirements, including the CHP, control worker overexposure to hazardous laboratory chemicals, thereby preventing serious illnesses and death among workers exposed to such chemicals.

**II. Special Issues for Comment**

OSHA has a particular interest in comments on the following issues:

- Whether the proposed information collection requirements are necessary for the proper performance of the Agency's functions to protect workers, including whether the information is useful;
- The accuracy of OSHA's estimate of the burden (time and costs) of the information collection requirements, including the validity of the methodology and assumptions used;
- The quality, utility, and clarity of the information collected; and
- Ways to minimize the burden on employers who must comply; for example, by using automated or other technological information collection and transmission techniques.

**III. Proposed Actions**

OSHA is proposing to extend the information collection requirements contained in the Occupational Exposure to Hazardous Chemicals in Laboratories Standard (29 CFR 1910.1450). The Agency is requesting to increase the existing burden hour estimate for the collection of information requirements in the Standard. In this regard, the Agency is requesting to adjust the current burden hour estimate from 281,419 hours to 293,706 hours.

*Type of Review:* Extension of currently approved collections.

*Title:* Occupational Exposure to Hazardous Chemicals in Laboratories.

*OMB Number:* 1218-0131.

*Affected Public:* Business or other for-profits.

*Number of Respondents:* 48,461.

*Frequency:* Varies from 3 minutes (.05 hour) to replace the safe practice manual to 1 hour to develop a new manual.

Total Responses: 948,634.

Average Time per Response:  
Annually; monthly, quarterly, semi-  
annually, on occasion

Estimated Total Burden Hours:  
293,706

Estimated Cost (Operation and  
Maintenance): \$41,271,276.

#### IV. Public Participation—Submission of Comments on This Notice and Internet Access to Comments and Submissions

You may submit comments in  
response to this document as follows:

(1) Electronically at <http://www.regulations.gov>, which is the Federal eRulemaking Portal; (2) by facsimile; or (3) by hard copy. All comments, attachments, and other material must identify the Agency name and the OSHA docket number for this ICR (Docket No. OSHA–2011–0059). You may supplement electronic submissions by uploading document files electronically. If you wish to mail additional materials in reference to an electronic or a facsimile submission, you must submit them to the OSHA Docket Office (see the section of this notice titled **ADDRESSES**). The additional materials must clearly identify your electronic comments by your name, date, and docket number so the Agency can attach them to your comments.

Because of security procedures, the use of regular mail may cause a significant delay in the receipt of comments. For information about security procedures concerning the delivery of materials by hand, express delivery, messenger or courier service, please contact the OSHA Docket Office at (202) 693–2350 (TTY (877) 889–5627).

Comments and submissions are posted without change at <http://www.regulations.gov>. Therefore, OSHA cautions commenters about submitting personal information such as Social Security Numbers and date of birth. Although all submissions are listed in the <http://www.regulations.gov> index, some information (e.g., copyrighted material) is not publicly available to read or download through this Web site. All submissions, including copyrighted material, are available for inspection and copying at the OSHA Docket Office. Information on using the <http://www.regulations.gov> Web site to submit comments and access the docket is available at the Web site's "User Tips" link. Contact the OSHA Docket Office for information about materials not available through the Web site, and for assistance in using the Internet to locate docket submissions.

#### V. Authority and Signature

David Michaels, PhD, MPH, Assistant Secretary of Labor for Occupational Safety and Health, directed the preparation of this notice. The authority for this notice is the Paperwork Reduction Act of 1995 (44 U.S.C. 3506 *et seq.*) and Secretary of Labor's Order No. 5–2010 (75 FR 55355).

Signed at Washington, DC, on April 28, 2011.

David Michaels,

Assistant Secretary of Labor for Occupational  
Safety and Health.

[FR Doc. 2011–10810 Filed 5–3–11; 8:45 am]

BILLING CODE 4510–26–P

#### THE NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

##### Federal Council on the Arts and the Humanities

##### Arts and Artifacts Indemnity Panel Advisory; Committee

**AGENCY:** The National Endowment for  
the Humanities.

**ACTION:** Notice of meeting.

Pursuant to the provisions of the Federal Advisory Committee Act (Pub. L. 92–463 as amended) notice is hereby given that a meeting of the Arts and Artifacts Indemnity Panel of the Federal Council on the Arts and the Humanities will be held at 1100 Pennsylvania Avenue, NW., Washington, DC 20506, in Room 730, from 9:30 a.m. to 5 p.m., on Thursday, May 26, 2011.

The purpose of the meeting is to review applications for Certificates of Indemnity submitted to the Federal Council on the Arts and the Humanities for exhibitions beginning after July 1, 2011.

Because the proposed meeting will consider financial and commercial data and because it is important to keep values of objects, methods of transportation and security measures confidential, pursuant to the authority granted me by the Chairman's Delegation of Authority to Close Advisory Committee Meetings, dated July 19, 1993, I have determined that the meeting would fall within exemption (4) of 5 U.S.C. 552(b) and that it is essential to close the meeting to protect the free exchange of views and to avoid interference with the operations of the Committee.

It is suggested that those desiring more specific information contact Advisory Committee Management Officer, Michael McDonald, 1100 Pennsylvania Avenue, NW.,

Washington, DC 20506, or call 202/606–8322.

Michael P. McDonald,

Management Officer, Advisory Committee.

[FR Doc. 2011–10789 Filed 5–3–11; 8:45 am]

BILLING CODE 7536–01–P

#### NATIONAL LABOR RELATIONS BOARD

##### Sunshine Act Meetings: May 2011

**TIME AND DATES:** All meetings are held at 2:30 p.m.

Tuesday, May 3; Wednesday, May 4; Thursday, May 5; Tuesday, May 10; Wednesday, May 11; Thursday, May 12; Tuesday, May 17; Wednesday, May 18; Thursday, May 19; Tuesday, May 24; Wednesday, May 25; Thursday, May 26; Tuesday, May 31.

**PLACE:** Board Agenda Room, No. 11820, 1099 14th St., NW., Washington DC 20570.

**STATUS:** Closed.

**MATTERS TO BE CONSIDERED:** Pursuant to § 102.139(a) of the Board's Rules and Regulations, the Board or a panel thereof will consider "the issuance of a subpoena, the Board's participation in a civil action or proceeding or an arbitration, or the initiation, conduct, or disposition \* \* \* of particular representation or unfair labor practice proceedings under section 8, 9, or 10 of the [National Labor Relations] Act, or any court proceedings collateral or ancillary thereto." See also 5 U.S.C. 552b(c)(10).

**CONTACT PERSON FOR MORE INFORMATION:**  
Lester A. Heltzer, Executive Secretary,  
(202) 273–1067.

Dated: May 2, 2011.

Lester A. Heltzer,

Executive Secretary.

[FR Doc. 2011–10992 Filed 5–2–11; 4:15 pm]

BILLING CODE 7545–01–P

#### NUCLEAR REGULATORY COMMISSION

[Docket Nos. 50–277 and 50–278; NRC–  
2011–0101]

##### Exelon Generation Company, LLC; PSEG Nuclear, LLC; Peach Bottom Atomic Power Station, Units 2 and 3; Notice of Withdrawal of Application for Amendment to Facility Operating License

The U.S. Nuclear Regulatory Commission (NRC or the Commission) has granted the request of Exelon Generation Company, LLC, (Exelon) and