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.

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

By Notice dated October 8, 2010, and published in the Federal Register on October 20, 2010, 75 FR 64745, 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:

<table>
<thead>
<tr>
<th>Drug</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dihydromorphine (9145)</td>
<td>I</td>
</tr>
<tr>
<td>Hydromorphone (9150)</td>
<td>II</td>
</tr>
</tbody>
</table>

Dihydromorphine is an intermediate in the manufacture of Hydromorphone and is not for commercial distribution. The company plans to manufacture Hydromorphone HCl for sale to other manufacturers and for the manufacture of other controlled substance dosage units 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: February 16, 2011.
Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 8, 2010, and published in the Federal Register on October 20, 2010, 75 FR 64745, 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:

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

DEPARTMENT OF JUSTICE

Parole Commission

[5 U.S.C. Section 552b]

Meetings: Sunshine Act; Public Announcement Pursuant to the Government in the Sunshine Act

AGENCY HOLDING MEETING: Department of Justice, United States Parole Commission.

DATE AND TIME: 10 a.m., Thursday, February 17, 2011.
PLACE: U.S. Parole Commission, 5550 Friendship Boulevard, 4th Floor, Chevy Chase, Maryland 20815.
STATUS: Closed.

MATTERS CONSIDERED: The following matter will be considered during the closed meeting: Discussion of an original jurisdiction case pursuant to 28 CFR 2.17.

AGENCY CONTACT: Patricia W. Moore, Staff Assistant to the Chairman, United States Parole Commission, (301) 492–5933.

Rockne Chickinell,
General Counsel, U.S. Parole Commission.

DEPARTMENT OF JUSTICE

United States Parole Commission

Record of Vote of Meeting Closure

(Pub. L. 94–409) (5 U.S.C. Sec. 552b)

I, Isaac Fulwood, of the United States Parole Commission, was present at a meeting of said Commission, which started at approximately 11:30 a.m., on Thursday, February 10, 2011, at the U.S. Parole Commission, 5550 Friendship Boulevard, 4th Floor, Chevy Chase, Maryland 20815. The purpose of the meeting was to decide four petitions for reconsideration pursuant to 28 CFR Section 2.27 and one pursuant to 28 CFR Section 2.17. Four Commissioners were present, constituting a quorum when the vote to close the meeting was submitted.

Public announcement further describing the subject matter of the meeting and certifications of General Counsel that this meeting may be closed by vote of the Commissioners present were submitted to the Commissioners prior to the conduct of any other business. Upon motion duly made, seconded, and carried, the following Commissioners voted that the meeting be closed: Isaac Fulwood, Cranston J. Mitchell, Patricia Cushwa and J. Patricia Wilson Smoot.

In witness whereof, I make this official record of the vote taken to close this meeting and authorize this record to be made available to the public.

Isaac Fulwood,
Chairman, U.S. Parole Commission.

DEPARTMENT OF LABOR

Office of Workers’ Compensation Programs

Division of Coal Mine Workers’ Compensation; Proposed Extension of Existing Collection; Comment Request

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a pre-clearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA95) [44 U.S.C. 3506(c)(2)(A)]. This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. Currently, the Office of Workers’ Compensation Programs is soliciting comments concerning its proposal to extend OMB approval of the information collection for the following medical reports:

<table>
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<th>Schedule</th>
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<td>Dihydromorphine (9145)</td>
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Roentgenographic Interpretation (CM–933), Roentgenographic Quality Rereading (CM–933b), Medical History and Examination for Coal Mine Workers’ Pneumoconiosis (CM–988), Report of Arterial Blood Gas Study (CM–1159) and Report of Ventilatory Study (CM–2907). A copy of the proposed information collection request can be obtained by contacting the office listed below in the addresses section of this Notice.

DATES: Written comments must be submitted to the office listed in the addresses section below on or before April 25, 2011.

ADDRESSES: Mr. Vincent Alvarez, U.S. Department of Labor, 200 Constitution Ave., NW., Room S–3201, Washington, DC 20210, telephone (202) 693–0372, fax (202) 693–1447, E-mail Alvarez.Vincent@ dol.gov. Please use only one method of transmission for comments (mail, fax, or E-mail).

SUPPLEMENTARY INFORMATION:

I. Background: The Black Lung Benefits Act of 1977 as amended, 20 U.S.C. 901 et seq. and 20 CFR 718.102 set forth criteria for the administration and interpretation of x-rays. When a miner applies for benefits, the Division of Coal Mine Workers’ Compensation (DCMWC) is required to schedule a series of four diagnostic tests to help establish eligibility for black lung benefits. Each of the diagnostic tests has its own form that sets forth the medical results. The forms are: Roentgenographic Interpretation (CM–933), Roentgenographic Quality Rereading (CM–933b), Medical History and Examination for Coal Mine Workers’ Pneumoconiosis (CM–988), Report of Arterial Blood Gas Study (CM–1159), and Report of Ventilatory Study (CM–2907). This information collection is currently approved for use through August 31, 2011.

II. Review Focus: The Department of Labor is particularly interested in comments which:

* 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

III. Current Actions: The Department of Labor seeks the approval for the extension of this currently-approved information collection in order to carry out its responsibility to administer the Black Lung Benefits Act.

Agency: Office of Workers’ Compensation Programs.

Title: Roentgenographic Interpretation (CM–933), Roentgenographic Quality Rereading (CM–933b), Medical History and Examination for Coal Mine Workers’ Pneumoconiosis (CM–988), Report of Arterial Blood Gas Study (CM–1159), and Report of Ventilatory Study (CM–2907).

OMB Number: 1240–0023.


Affected Public: Business or other for profit, and not-for-profit institutions.

<table>
<thead>
<tr>
<th>Form</th>
<th>Time to complete</th>
<th>Frequency of response</th>
<th>Number of respondents</th>
<th>Number of responses</th>
<th>Hours burden</th>
</tr>
</thead>
<tbody>
<tr>
<td>CM–933</td>
<td>5 min</td>
<td>on occasion</td>
<td>4800</td>
<td>4800</td>
<td>400</td>
</tr>
<tr>
<td>CM–933b</td>
<td>3 min</td>
<td>on occasion</td>
<td>4800</td>
<td>4800</td>
<td>240</td>
</tr>
<tr>
<td>CM–988</td>
<td>30 min</td>
<td>on occasion</td>
<td>4800</td>
<td>4800</td>
<td>2400</td>
</tr>
<tr>
<td>CM–1159</td>
<td>15 min</td>
<td>on occasion</td>
<td>4800</td>
<td>4800</td>
<td>1200</td>
</tr>
<tr>
<td>CM–2907</td>
<td>20 min</td>
<td>on occasion</td>
<td>4800</td>
<td>4800</td>
<td>1600</td>
</tr>
<tr>
<td>Totals</td>
<td></td>
<td></td>
<td>24000</td>
<td>24000</td>
<td>5840</td>
</tr>
</tbody>
</table>

Total Respondents: 24,000.
Total Annual Responses: 24,000.
Average Time per Response: 3 minutes–30 minutes.
Estimated Total Burden Hours: 5,840.
Frequency: On occasion.
Total Burden Cost (capital/startup): $50.
Total Burden Cost (operating/maintenance): $35,520.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget approval of the information collection request; they will also become a matter of public record.

Dated: February 16, 2011.

Vincent Alvarez,
Agency Clearance Officer, Office of Workers’ Compensation Programs, U.S. Department of Labor.

DEPARTMENT OF LABOR
Office of Workers’ Compensation Programs

Division of Federal Employees’ Compensation; Proposed Extension of Existing Collection; Comment Request

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995 [PRA95] [44 U.S.C. 3506(c)(2)(A)]. This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. Currently, the Office of Workers’ Compensation Programs is soliciting comments concerning the proposed collection: Notice of Recurrences (CA–2a). A copy of the proposed information collection request can be obtained by contacting the office listed below in the addresses section of this Notice.

DATES: Written comments must be submitted to the office listed in the addresses section below on or before April 25, 2011.

ADDRESSES: Mr. Vincent Alvarez, U.S. Department of Labor, 200 Constitution Ave., NW., Room S–3201, Washington, DC 20210, telephone (202) 693–0372, fax (202) 693–1447, E-mail