Metallic Mineral Processing Plants (40 CFR part 60, subpart LL) were promulgated on February 21, 1984, and amended on October 17, 2000. This NSPS affects both owners and operators of metallic mineral processing plants.

Owners and operators must conduct initial performance tests, maintain records of startups, shutdowns, malfunctions, and continuous monitoring system parameters, and submit semianual reports. These notifications, reports, and records are essential in determining compliance; and, in general, are required of all sources subject to NSPS.

Notifications are to inform the Agency or delegated authority when a source becomes subject to the standard. The reviewing authority may then inspect the source to check if the pollution control devices are properly installed and operating, and that the standards are being met. Performance test reports are required as these are the Agency’s records of a source’s initial capability to comply with the emission standards and to serve as a record of the operating conditions under which compliance are to be achieved. The information generated by monitoring, recordkeeping, and reporting requirements described in this ICR are used by the Agency to ensure that facilities affected by the standard continue to operate the control equipment and achieve continuous compliance with the regulation.

All reports are sent to the delegated state or local authority. In the event that there is no such delegated authority, the reports are sent directly to the EPA regional office. This information is being collected to assure compliance with 40 CFR part 60, subpart LL, as authorized in section 112 and 114(a) of the Clean Air Act. The required information consists of emissions data and other information that have been determined to be private.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB Control Number. The OMB Control Numbers for the EPA regulations are listed in 40 CFR part 9 and 48 CFR chapter 15, and are identified on the form and/or instrument, if applicable.

Burden Statement: The annual public reporting and recordkeeping burden for this collection of information is estimated to average 52 hours per response. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, or disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; develop, acquire, install, and utilize technology and systems for the purposes of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; adjust the existing ways to comply with any previously applicable instructions and requirements which have subsequently changed; train personnel to be able to respond to a collection of information; search data sources; complete and review the collection of information; and transmit or otherwise disclose the information.

Respondents/Affected Entities: Metallic mineral processing plants.

Estimated Number of Respondents: 20.

Frequency of Response: Initially, occasionally, and semianually.

Estimated Total Annual Hour Burden: 2,306.

Estimated Total Annual Cost: $233,712, which includes $220,712 in labor costs, no capital/startup costs, and $13,000 in operation and maintenance (O&M) costs.

Changes in the Estimates: There is no change in the labor hours to the respondents in this ICR compared to the previous ICR. After consulting the Office of Air Quality Planning and Standards (OAQPS) and trade associations, our data indicates that there are approximately 20 sources subject to the rule, with no additional new sources over the next three years.

However, there is an increase in the estimated burden cost as currently identified in the OMB Inventory of approved Burdens. The increase is not due to any program changes, but the change in burden is due to the use of the most updated labor rates.

John Moses,
Director, Collection Strategies Division.

SUPPLEMENTARY INFORMATION:
I. General Information
A. Does this action apply to me?

This action is directed to the public in general, and may be of interest to a wide range of stakeholders including environmental, human health, and agricultural advocates; the chemical industry; pesticide users; and members of the public interested in the sale, distribution, or use of pesticides. Since others also may be interested, the Agency has not attempted to describe all the specific entities that may be affected by this action. If you have any questions regarding the applicability of this action to a particular entity, consult the person listed under FOR FURTHER INFORMATION CONTACT.

B. How can I get copies of this document and other related information?

EPA has established a docket for this action under docket identification (ID) number EPA–HQ–OPP–2005–0220. Publicly available docket materials are available either in the electronic docket at http://www.regulations.gov, or, if only...
available in hard copy, at the Office of Pesticide Programs (OPP) Regulatory Public Docket in Rm. S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Dr. Arlington, VA. The hours of operation of this Docket Facility are from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The Docket Facility telephone number is (703) 305–5805.

II. What action is the agency taking?

This notice announces the cancellation of dicofol products, as requested by registrants, pursuant to section 6(f) of FIFRA. These registrations are listed in sequence by EPA company number. Tables 1 and 2 of this unit, in sequence by EPA company number, list the names and addresses of record for all registrations are listed in sequence by section 6(f) of FIFRA. These registrations are requested by registrants, pursuant to cancellation of dicofol products, as effective date of the action. The existing registrations identified in Tables 1 and 2 of Unit II. Accordingly, the cancellation of the technical dicofol product is effective immediately. The cancellation of the end use registrations is effective October 31, 2013. Any distribution, sale, or use of existing stocks of the products identified in Tables 1 and 2 of Unit II in a manner inconsistent with any of the provisions for disposition of existing stocks set forth in Unit VI will be a violation of FIFRA.

III. Summary of Public Comments Received and Agency Response to Comments

During the public comment period provided, EPA received no comments in response to the June 22, 2011 Federal Register notice announcing the Agency’s receipt of the requests for voluntary cancellations of products listed in Tables 1 and 2 of Unit II.

IV. Cancellation Order

Pursuant to FIFRA section 6(f), EPA hereby approves the requested cancellations and amendments to terminate the uses of dicofol registrations identified in Tables 1 and 2 of Unit II. Accordingly, the cancellation of the technical dicofol product is effective immediately. The cancellation of the end use registrations is effective October 31, 2013. Any distribution, sale, or use of existing stocks of the products identified in Tables 1 and 2 of Unit II in a manner inconsistent with any of the provisions for disposition of existing stocks set forth in Unit VI will be a violation of FIFRA.

V. What is the agency’s authority for taking this action?

Section 6(f)(1) of FIFRA provides that a registrant of a pesticide product may at any time request that any of its pesticide registrations be canceled or amended to terminate one or more uses. FIFRA further provides that, before acting on the request, EPA must publish a notice of receipt of any such request in the Federal Register. Thereafter, following the public comment period, the EPA Administrator may approve such a request. The notice of receipt for this action was published for comment on June 22, 2011 (76 FR 36535) (FRL–8875–7). The comment period closed on July 22, 2011.

VI. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products which are currently in the United States and which were packaged, labeled, and released for shipment prior to the effective date of the action. The existing stocks provision for the products subject to this order is as follows:

a. Sale, distribution and use of the technical dicofol is prohibited, except: Registrants of dicofol end-use products shall be allowed to reformulate existing stocks of dicofol technical into products identified in Table 2 of Unit II, until October 31, 2013, or for products intended for export consistent with the requirements of FIFRA section 17 or for purposes of proper disposal.

b. Sale and distribution by registrants of end use products after October 31, 2013 is prohibited except for export consistent with the requirements of FIFRA section 17 or for purposes of proper disposal.

c. Sale and distribution of end use products by persons other than registrants is permitted until December 31, 2013. Thereafter, sale and distribution of end use products by persons other than registrants is prohibited except for export consistent with the requirements of FIFRA section 17 or for purposes of proper disposal.

d. Use of existing stocks of any end-use product consistent with the terms of the previously approved labeling on, or accompanied by, the deleted products identified shall be allowed until October 31, 2016, and thereafter, only for the purposes of proper disposal.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: December 6, 2011.

Richard P. Keigwin, Jr.,
Director, Pesticide Re-evaluation Division,
Office of Pesticide Programs.

[FR Doc. 2011–31987 Filed 12–13–11; 8:45 am]
BILLING CODE 6550–50–P

ENVIRONMENTAL PROTECTION AGENCY

[76 FR 77825]

Human Studies Review Board;
Notification of a Public Teleconference

AGENCY: Environmental Protection Agency.

ACTION: Notice.

SUMMARY: The EPA Office of the Science Advisor announces a public teleconference of the HSRB to discuss its draft report from the October 19–20, 2011 HSRB meeting.

DATES: The teleconference will be held on Wednesday, January 11, 2012 from approximately 1 p.m. to approximately 4 p.m. Eastern Time. Comments may be submitted on or before Wednesday, January 4, 2012.

ADDRESSES: Submit your written comments, identified by Docket ID No. EPA–HQ–ORD–2011–0953, by one of the following methods:


Email: ORD.Docket@epa.gov.


Hand Delivery: The EPA/DC Public Reading Room is located in the EPA Headquarters Library, Room Number 3334 in the EPA West Building, located at 1301 Constitution Avenue NW, Washington, DC 20460. The hours of operation are 8:30 a.m. to 4:30 p.m.

Table 1—Dicofol Technical Product Cancellation

<table>
<thead>
<tr>
<th>EPA registration No.</th>
<th>Product name</th>
</tr>
</thead>
<tbody>
<tr>
<td>11603–26</td>
<td>Mitigan (Dicofol) Technical</td>
</tr>
</tbody>
</table>

Table 2—Dicofol End Use Product Cancellations

<table>
<thead>
<tr>
<th>EPA registration No.</th>
<th>Product name</th>
</tr>
</thead>
<tbody>
<tr>
<td>66222–21</td>
<td>MANA Dicofol 4E</td>
</tr>
<tr>
<td>66222–56</td>
<td>Dicofol 4E</td>
</tr>
<tr>
<td>66222–95</td>
<td>Dicofol 50WSB</td>
</tr>
</tbody>
</table>

Table 3 of this unit includes the names and addresses of record for all registrants of the products listed in Tables 1 and 2 of this unit, in sequence by EPA company number.

Table 3—Registrants of Cancelled Products

<table>
<thead>
<tr>
<th>EPA company No.</th>
<th>Company name and address</th>
</tr>
</thead>
<tbody>
<tr>
<td>11603</td>
<td>Agan Chemical Manufacturing Ltd, 4515 Falls of Neuse Rd, Suite 300, Raleigh, NC 27609.</td>
</tr>
</tbody>
</table>