to leak detection and repair (LDAR) and pressure relief devices (PRD) for subject PC and AMF facilities. We assume existing PC and AMF facilities will come into compliance with the new requirements during the three-year period covered under this ICR.

Courtney Kerwin,

Acting Director, Collection Strategies Division.

[FR Doc. 2015–18659 Filed 7–29–15; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OECA–2014–0025; FRL—9931–57–OEI]

Information Collection Request Submitted to OMB for Review and Approval; Comment Request; NESHAP for Asbestos (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency has submitted an information collection request (ICR), ''NESHAP for Asbestos (40 CFR part 61, subpart M) (Renewal)'' (EPA ICR No. 0111.14, OMB Control No. 2060–0101) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act (44 U.S.C. 3501 *et seq.*). This is a proposed extension of the ICR, which is currently approved through July 31, 2015. Public comments were previously requested via the **Federal Register** (79 FR 30117) on May 27, 2014 during a 60-day comment period. This notice allows for an additional 30 days for public comments. A fuller description of the ICR is given below, including its estimated burden and cost to the public. 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.

DATES: Additional comments may be submitted on or before August 31, 2015.

ADDRESSES: Submit your comments, referencing Docket ID Number EPA–HQ–OECA–2014–0025, to (1) EPA online using *www.regulations.gov* (our preferred method), by email to *docket.oeca@epa.gov,* or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW., Washington, DC 20460, and (2) OMB via email to *oira_submission@omb.eop.gov.* Address comments to OMB Desk Officer for EPA.

EPA's policy is that all comments received will be included in the public docket without change including any personal information provided, unless the comment includes profanity, threats, information claimed to be Confidential Business Information (CBI), or other information whose disclosure is restricted by statute.

FOR FURTHER INFORMATION CONTACT: Patrick Yellin, Monitoring, Assistance, and Media Programs Division, Office of Compliance, Mail Code 2227A, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460; telephone number: (202) 564–2970; fax number: (202) 564–0050; email address: *yellin.patrick@epa.gov.*

SUPPLEMENTARY INFORMATION: Supporting documents which explain in detail the information that the EPA will be collecting are available in the public docket for this ICR. The docket can be viewed online at *www.regulations.gov* or in person at the EPA Docket Center, EPA West, Room 3334, 1301 Constitution Ave. NW., Washington, DC. The telephone number for the Docket Center is (202) 566–1744. For additional information about EPA's public docket, visit *www.epa.gov/dockets.*

Abstract: The affected entities are subject to the General Provisions of the NESHAP at 40 CFR part 61, subpart A, and any changes, or additions to the General Provisions specified at 40 CFR part 61, subpart M. Owners or operators of the affected facilities must submit initial notification, performance tests, and periodic reports and results. Owners or operators are also required to maintain records of the occurrence and duration of any startup, shutdown, or malfunction in the operation of an affected facility, or any period during which the monitoring system is inoperative. Reports, at a minimum, are required semiannually.

Form Numbers: None.

Respondents/affected entities: Demolition and renovation facilities; disposal of asbestos wastes; asbestos milling, manufacturing and fabricating; use of asbestos on roadways; asbestos waste conversion facilities; and the use of asbestos insulation and spray-on materials.

Respondent's obligation to respond: Mandatory (40 CFR part 61, subpart M).

Estimated number of respondents: 9,603 (total).

Frequency of response: Initially, occasionally, quarterly and semiannually.

Total estimated burden: 292,050 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated cost: $29,370,000 (per year), includes $0 annualized capital or operation & maintenance costs.

Changes in the Estimates: There is an increase in the total estimated burden as currently identified in the OMB Inventory of Approved Burdens. The change is due to an increase in the asbestos demolition and renovation operations each year; it is not due to any program changes. We have updated respondent and Agency burdens to reflect EPA's current estimates of sources subject to the rule. We have also updated respondent and Agency labor rates, which were referenced from the Bureau of Labor Statistics and OPM, respectively.

During the preparation of this ICR, EPA identified several discrepancies related to rule reporting/recordkeeping requirements and associated burdens. We have updated the respondent and Agency burden tables accordingly in order to reconcile the discrepancies. For example, the previous ICR did not correctly reflect the number of demolition/renovation contractors that will participate in refresher training. The revisions did not result in a substantial burden change, as the average reporting and recordkeeping burden hours per response in this ICR is equal to that of the previous ICR.

Courtney Kerwin,

Acting Director, Collection Strategies Division.

[FR Doc. 2015–18660 Filed 7–29–15; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[CERCLA–04–2015–3752; FRL–9931–52–Region 4]

Capitol City Plume Superfund Site Montgomery, Montgomery County, Alabama; Notice of settlement

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of settlement.

SUMMARY: Under 122(h) of the Comprehensive Environmental Response, Compensation and Liability Act (CERCLA), the United States Environmental Protection Agency (EPA) has entered into a settlement with multiple parties concerning the Capitol City Plume Superfund Site located in Montgomery, Montgomery County, Alabama. The settlement addresses costs from a fund-lead Remedial Investigation performed by EPA at the Site. The Agency is deferring the Site to the State of Alabama for cleanup.

DATES: The Agency will consider public comments on the settlement until August 31, 2015. The Agency will consider all comments received and may modify or withdraw its consent to the settlement if comments received disclose facts or considerations which indicate that the amended settlement is inappropriate, improper, or inadequate.

ADDRESSES: Copies of the settlement are available from the Agency by contacting Ms. Paula V. Painter, Environmental Protection Specialist using the contact information provided in this notice. Comments may also be submitted by referencing the Site's name through one of the following methods:

- *Internet: www.epa.gov/region4/superfund/programs/enforcement/enforcement.html.*
- *U.S. Mail:* U.S. Environmental Protection Agency, Superfund Division, Attn: Paula V. Painter, 61 Forsyth Street SW., Atlanta, Georgia 30303.
- *Email: Painter.Paula@epa.gov.*

FOR FURTHER INFORMATION CONTACT: Paula V. Painter at (404) 562–8887.

Dated: June 9, 2015.

Anita L. Davis,

Chief, Enforcement and Community Engagement Branch, Superfund Division.

[FR Doc. 2015–18727 Filed 7–29–15; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0397]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; State Enforcement Notifications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled, ''State Enforcement Notifications'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On June 8, 2015, the Agency submitted a proposed collection of information entitled, ''State Enforcement Notifications'' to OMB for review and clearance under 44 U.S.C. 3507. 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. OMB has now approved the information collection and has assigned OMB control number 0910–0275. The approval expires on July 31, 2018. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: July 24, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–18649 Filed 7–29–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–N–1305]

Multicriteria-Based Ranking Model for Risk Management of Animal Drug Residues in Milk and Milk Products; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or we) is extending the comment period for the risk assessment entitled ''Multicriteria-Based Ranking Model for Risk Management of Animal Drug Residues in Milk and Milk Products.'' A notice of the availability of the risk assessment and our request for comments appeared in the **Federal Register** of April 30, 2015. We initially established July 29, 2015, as the deadline for the submission of requested comments that can help improve the ranking model approach, including the specific criteria, scoring, and weighting scheme; the scientific data and assumptions used to inform scoring used in the model; the selection of animal drugs evaluated; and the clarity and the transparency of the risk assessment. We are taking this action in response to a request for an extension to allow interested persons additional time to submit comments.

DATES: FDA is extending the comment period on the risk assessment whose availability we announced in a notice published on April 30, 2015 (80 FR 24260). Submit either electronic or written comments on the risk assessment by October 27, 2015.

ADDRESSES: You may submit comments by any of the following methods:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.

Written Submissions

Submit written submissions in the following ways:

- *Mail/Hand delivery/Courier (for paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Instructions: All submissions received must include the Docket No. FDA–2015–N–1305. All comments received may be posted without change to *http://www.regulations.gov,* including any personal information provided. For additional information on submitting comments, see the ''Comments'' heading of the **SUPPLEMENTARY INFORMATION** section of this document.

Docket: For access to the docket to read background documents or comments received, go to *http://www.regulations.gov* and insert the docket number(s), found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Jane Van Doren, Center for Food Safety and Applied Nutrition (HFS–005), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 240–402–2927.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of April 30, 2015, FDA published a notice announcing the availability of a risk assessment entitled ''Multicriteria-Based Ranking Model for Risk Management of Animal Drug Residues in Milk and Milk Products,'' with a 90-day comment period to request comments on the risk assessment.

We received a request for a 90-day extension of the comment period for the risk assessment. The request conveyed concern that the current 90-day comment period does not allow sufficient time to develop meaningful or thoughtful comments to the risk assessment.

FDA has considered the request and is extending the comment period for the risk assessment for 90 days, until October 27, 2015. We believe that a 90-