final registration review decision for soap salts will depend upon the results of an ESA section 7 consultation with the services, an EDSP determination, and an assessment of the non-target exposure to bees.

Sulfur (Interim Decision). The registration review docket for sulfur (EPA–HQ–OPP–2008–0176) opened in March 2008. Sulfur is used as an insecticide and fungicide on a wide range of field and greenhouse-grown food and feed crops, livestock, livestock quarters, and indoor and outdoor residential sites. Sulfur is also registered for use in gas cartridge products, along with inorganic nitrate/nitrite, carbon, and carbon dioxide. EPA has conducted a qualitative assessment for both human health and ecological risks, including listed species, for sulfur. Details of the assessment for the gas cartridge use are summarized under the gas cartridge heading in this unit. For uses of sulfur other than gas cartridges, the Agency is making a ''no effect'' determination for all listed aquatic species, and a ''no effect'' determination for direct effects to listed terrestrial vertebrates that do not rely on insects as a primary food source. However, at this time, the Agency is not able to make a listed species determination for effects to terrestrial invertebrates, terrestrial plants, or indirect effects to terrestrial vertebrates with insects as a primary food source. Sulfur has not been evaluated under the EDSP. Therefore, the Agency's final registration review decision is dependent upon the result of an ESA Section 7 consultation with the USFWS and the evaluation of potential endocrine disruptor risk.

Case Closure for Flufenpyr-ethyl (PC Code 108853; Case 7262). Flufenpyr-ethyl is an herbicide which was labeled for post-emergence control of broadleaf weeds in field corn, soybeans, and sugarcane. On March 19, 2015, the Agency received a request for voluntary cancellation of flufenpyr-ethyl from the technical and end-use product registrant, Valent USA Corporation. EPA subsequently issued a **Federal Register** notice announcing receipt of the request (FRL–9928–54) on July 8, 2015 (80 FR 39100), and allowed for a 30-day period for public comment on the request. No comments were received, and on September 22, 2015, EPA issued the cancellation order terminating the last pesticide products containing flufenpyr-ethyl registered in the United States (80 FR 57179) (FRL–9933–58). There were no existing stocks of these products and no requests for existing stocks provisions. Therefore no existing stocks provision was provided for these product registrations. With the cancellation of these remaining products, the Agency is announcing the closure of the registration review case for flufenpyr-ethyl.

Pursuant to 40 CFR 155.57, a registration review decision is the Agency's determination whether a pesticide meets, or does not meet, the standard for registration in FIFRA. EPA has considered the pesticides listed in light of the FIFRA standard for registration. The interim decision documents in the dockets describe the Agency's rationale for issuing registration review interim decisions for these pesticides.

In addition to the interim registration review decision documents, the registration review docket for these pesticides also includes other relevant documents related to the registration review of these cases. The proposed interim registration review decisions were previously posted to each docket and the public was invited to submit any comments or new information.

EPA has addressed the substantive comments or information received during the 60-day comment period in the interim decision document for each pesticide listed in this document.

Pursuant to 40 CFR 155.58(c), the registration review case docket for each pesticide discussed in this notice will remain open until all actions required in the interim decisions have been completed.

Background on the registration review program is provided at: *http://www.epa.gov/oppsrrd1/registration_review.* Links to earlier documents related to the registration review of this pesticide are provided in the Pesticide Chemical Search data base accessible at: *http://iaspub.epa.gov/apex/pesticides/f?p=chemicalsearch.*

Authority: 7 U.S.C. 136 *et seq.*

Dated: September 30, 2015.

Bernard P. Keigwin, Jr.,

Director, Pesticide Re-Evaluation Division, Office of Pesticide Programs.

[FR Doc. 2015–26299 Filed 10–14–15; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OAR–2003–0052; FRL–9935–45–OEI]

Information Collection Request Submitted to OMB for Review and Approval; Comment Request; Risk Management Program Requirements and Petitions To Modify the List of Regulated Substances (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency has submitted an information collection request (ICR), ''Risk Management Program Requirements and Petitions to Modify the List of Regulated Substances under section 112(r) of the Clean Air Act (CAA)'' (EPA ICR No. 1656.15, OMB Control No. 2050–0144) 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 December 31, 2015. Public comments were previously requested via the **Federal Register** (80 FR 33518) on June 12, 2015 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 November 16, 2015.

ADDRESSES: Submit your comments, referencing Docket ID No. EPA–HQ–OAR–2003–0052, to (1) EPA online using *www.regulations.gov* (our preferred method), 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: James Belke, Office of Emergency Management, Mail Code 5104A, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460; telephone number: 202–564–8023; fax number: 202–564–2625; email address: *belke.jim@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, WJC 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 *http://www.epa.gov/dockets.*

Abstract: The 1990 CAA Amendments added section 112(r) to provide for the prevention and mitigation of accidental releases. Section 112(r) mandates that EPA promulgate a list of ''regulated substances'' with threshold quantities and establish procedures for the addition and deletion of substances from the list of regulated substances. Processes at stationary sources that contain more than a threshold quantity of a regulated substance are subject to accidental release prevention regulations promulgated under CAA section 112(r)(7). These two rules are codified as 40 CFR part 68. Part 68 requires that sources with more than a threshold quantity of a regulated substance in a process develop and implement a risk management program and submit a risk management plan to EPA. The compliance schedule for the Part 68 requirements, established by rule on June 20, 1996, requires the implementation of the source risk management programs and the submission of initial Risk Management Plans (RMPs) by June 21, 1999, and at least every five years after the initial submission. Sources must resubmit earlier than their next five-year deadline if they undergo certain changes to their covered processes as specified in Part 68. Therefore, after the initial submission, some sources re-submitted their RMPs prior to the next 5-year deadline because they had process changes that required an earlier update. These sources were then assigned a new five-year resubmission deadline based on the date of their revised plan submission. Most covered sources had no significant changes to their covered processes and therefore resubmitted their updated RMP on June 21, 2004. This same pattern continued through the next two submission cycles—some sources updated and resubmitted their RMP prior to their next five-year deadline and were assigned a new (off-cycle) five-year deadline, but a majority of sources submitted their updated RMP on or near the next scheduled five-year resubmission deadlines (June 2009 and June 2014). Similarly, while most sources' next submission is due in June 2019, because of off-cycle resubmission deadlines assigned to sources who have resubmitted RMPs prior to their next 5-year resubmission date, only a portion of the RMP-regulated universe has a submission deadline occurring in June 2019.

Other than the costs for gathering information and filling out the on-line RMP form, the regulations require sources to maintain on-site documentation, perform a compliance audit every three years, provide refresher training to employees, perform a hazard analysis at least every five years, etc. Some of these activities are expected to occur annually or are on-going. Some are required every three years or every five years, unless there are changes at the facility. Therefore, the burden and costs incurred by sources vary from ICR to ICR. The five-year resubmission deadline set by the regulations or assigned by EPA based on the latest RMP resubmission also will cause the burden to vary from ICR to ICR.

Form Numbers: 8700–25, 8700–27, 8700–28.

Respondents/affected entities: Chemical manufacturers, petroleum refineries, water treatment systems, agricultural chemical distributors, refrigerated warehouses, chemical distributors, non-chemical manufacturers, wholesale fuel distributors, energy generation facilities, etc.

Respondent's obligation to respond: Mandatory (40 CFR part 68).

Estimated number of respondents: 13,396 (total).

Frequency of response: On occasion.

Total estimated burden: 54,000 hours (per year). Burden is defined at 5 CFR 1320.03(b)

Total estimated cost: $6,680,625 (per year), includes $0 annualized capital or operation & maintenance costs.

Changes in the estimates: There is a decrease of 26,546 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. The reason for this decrease is because this ICR period does not include a major filing deadline year and the previous ICR did include a major filing deadline. Second, the number of sources subject to the regulations fluctuates regularly, and is lower in this ICR period than in the previous ICR.

Courtney Kerwin,

Acting Director, Collection Strategies Division.

[FR Doc. 2015–26231 Filed 10–14–15; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL DEPOSIT INSURANCE CORPORATION

FDIC Advisory Committee on Economic Inclusion (ComE–IN); Notice of Meeting

AGENCY: Federal Deposit Insurance Corporation (FDIC).

ACTION: Notice of open meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, notice is hereby given of a meeting of the FDIC Advisory Committee on Economic Inclusion, which will be held in Washington, DC. The Advisory Committee will provide advice and recommendations on initiatives to expand access to banking services by underserved populations.

DATES: Friday, October 30, 2015, from 9 a.m. to 3:30 p.m.

ADDRESSES: The meeting will be held in the FDIC Board Room on the sixth floor of the FDIC Building located at 550 17th Street NW., Washington, DC.

FOR FURTHER INFORMATION CONTACT: Requests for further information concerning the meeting may be directed to Mr. Robert E. Feldman, Committee Management Officer of the FDIC, at (202) 898–7043.

SUPPLEMENTARY INFORMATION:

Agenda: The agenda will be focused on the Bank On 2.0 initiative, mobile banking research, expanding economic inclusion for individuals with disabilities, and Money Smart for Small Business. The agenda may be subject to change. Any changes to the agenda will be announced at the beginning of the meeting.

Type of Meeting: The meeting will be open to the public, limited only by the space available on a first-come, first-served basis. For security reasons, members of the public will be subject to security screening procedures and must present a valid photo identification to enter the building. The FDIC will provide attendees with auxiliary aids (*e.g.,* sign language interpretation) required for this meeting. Those attendees needing such assistance should call (703) 562–6067 (Voice or TTY) at least two days before the meeting to make necessary arrangements. Written statements may