

Dated: January 31, 2018,
Kimberly Y. Patrick,
Director, Office of Acquisition Management.
 [FR Doc. 2018-02929 Filed 2-12-18; 8:45 am]
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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OECA-2017-0640; FRL-9974-23-OECA]

Proposed Information Collection Request; Comment Request; Recordkeeping Requirements for Producers, Registrants and Applicants of Pesticides and Pesticide Devices Under Section 8 of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA); EPA ICR Number 0143.13, OMB Control Number 2070-0028

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency is planning to submit an information collection request (ICR), Recordkeeping Requirements for Producers, Registrants and Applicants of Pesticides and Pesticide Devices under Section 8 of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) (EPA ICR No. 0143.13, OMB Control No. 2070-0028) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act. EPA is soliciting public comments on specific aspects of the proposed information collection as described below. This is a proposed extension of the ICR, which is currently approved through September 30, 2018. 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: Comments must be submitted on or before April 16, 2018.

ADDRESSES: Submit your comments, referencing Docket ID No. EPA-HQ-OECA-2017-0640, 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.

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:

Michelle Stevenson, Office of Compliance, Monitoring, Assistance, and Media Programs Division, Pesticides, Waste & Toxics Branch (2225A), Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: (202) 564-4203; fax number: (202) 564-0085; email: stevenson.michelle@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>.

Pursuant to section 3506(c)(2)(A) of the PRA, EPA is soliciting comments and information to enable it to: (i) 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; (ii) 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; (iii) enhance the quality, utility, and clarity of the information to be collected; and (iv) minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses. EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval. At that time, EPA will issue another **Federal Register** notice to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB.

Abstract: Producers of pesticides and pesticide devices must maintain certain records with respect to their operations and make such records available for inspection and copying as specified in Section 8 of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and in regulations at 40 CFR part 169.

This information collection is mandatory under FIFRA Section 8. It is used by the Agency to determine compliance with FIFRA. The information is used by EPA Regional pesticide enforcement and compliance staffs, the Office of Enforcement and Compliance Assurance (OECA), and the Office of Pesticide Programs (OPP) within the Office of Chemical Safety and Pollution Prevention (OCSPP), as well as the U.S. Department of Agriculture (USDA), the Food and Drug Administration (FDA), and other Federal agencies, States under Cooperative Enforcement Agreements, and 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.

Form Numbers: None.

Respondents/affected entities:

Producers of pesticides and pesticide devices for sale or distribution in or exported to the United States.

Respondent's obligation to respond: Mandatory (40 CFR 169).

Estimated number of respondents: 14,447 (total).

Frequency of response: Annual.

Total estimated burden: 28,894

Total estimated cost: \$3,500,508.

There are no annualized capital or O&M costs associated with this ICR since all equipment associated with this ICR is present as part of ordinary business practices.

Changes in estimates: There is a decrease of 5,694 hours in the total estimated burden currently identified in the OMB Inventory of Approved ICR Burdens. This decrease is an adjustment due to a change in the number of respondents since the last ICR.

Dated: January 11, 2018.

Edward J. Messina,

Director, Office of Compliance/MAMPD.

[FR Doc. 2018-02931 Filed 2-12-18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of

the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP); Funding Opportunity Announcement (FOA), IP18–001, Reducing Disparities in Vaccination Coverage by Poverty Status Among Young Children and IP18–003, Understanding and Addressing the Disparity in Vaccination Coverage Among U.S. Adolescents Living in Rural versus Urban Areas.

Dates: March 20, 2018 and March 21, 2018.
Time: 10:00 a.m.–5:00 p.m. (EDT)

Place: Teleconference, Centers for Disease Control and Prevention, Room 1080, 8 Corporate, Atlanta, GA 30329.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Gregory Anderson, M.S., M.P.H., Scientific Review Officer, CDC, 1600 Clifton Road NE, Mailstop E–60, Atlanta, Georgia 30333, (404) 718–8833, gca5@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** Notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2018–02826 Filed 2–12–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Lead Exposure and Prevention Advisory Committee (LEPAC); Notice of Establishment

ACTION: Notice of charter establishment.

SUMMARY: Pursuant to the Section 2203 of Public Law 114–322 (Water Infrastructure Improvements for the Nation Act)(Registry for Lead Exposure and Advisory Committee), and the Federal Advisory Committee Act of October 6, 1972, the Director, Centers for Disease Control and Prevention (CDC), announces the establishment of the Lead Exposure and Prevention Advisory Committee. The Lead

Exposure and Prevention Advisory Committee shall, at a minimum: (1) Review the Federal programs and services available to individuals and communities exposed to lead; (2) review current research on lead exposure to identify additional research needs; (3) review and identify best practices, or the need for best practices regarding lead screening and the prevention of lead poisoning; (4) identify effective services, including services relating to healthcare, education, and nutrition for individuals and communities affected by lead exposure and lead poisoning, including in consultation with, as appropriate, the lead exposure registry as established in Section 2203(b) of Public Law 114–322; and (5) undertake any other review or activities that the Secretary determines to be appropriate. This advisory committee will review research and Federal programs and services related to lead poisoning and to identify effective services and best practices for addressing and preventing lead exposures in communities.

FOR FURTHER INFORMATION CONTACT:

Perri Ruckart, M.P.H., Epidemiologist, CDC, 4770 Buford Highway NE, Atlanta, Georgia 30341, telephone (770) 488–3808; afp4@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2018–02823 Filed 2–12–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

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The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463. The grant applications and

the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel; (SEP) DP18–001, Etiology and Outcome of Inflammatory Bowel Disease.

Date: April 11, 2018.

Time: 11:00 a.m.–6:00 p.m., EDT.

Place: Teleconference.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Jaya Raman Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway, Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488–6511, kva5@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2018–02829 Filed 2–12–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

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Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP); DD18–001, Birth Defects Study To Evaluate Pregnancy exposureS (BD–STEPS) II.