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ADDRESSES: Follow the detailed instructions provided under **ADDRESSES** in the **Federal Register** document of July 14, 2016 (81 FR 45477) (FRL-9948–29).

FOR FURTHER INFORMATION CONTACT:

Margaret Hathaway, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 305–5076; email address: hathaway.margaret@epa.gov.

SUPPLEMENTARY INFORMATION: This document reopens the public comment period established in the Federal Register document of July 14, 2016. In that document, EPA opened a public comment period for a proposed interim decision for 22 sulfonylurea herbicides: Bensulfuron-methyl, chlorimuron-ethyl, chlorsulfuron, flazasulfuron, foramsulfuron, halosulfuron-methyl, imazosulfuron, iodosulfuron-methylsodium, mesosulfuron-methyl, metsulfuron-methyl, nicosulfuron, orthosulfamuron, primisulfuron-methyl, prosulfuron, rimsulfuron, sulfometuronmethyl, sulfosulfuron, thifensulfuronmethyl, triasulfuron, tribenuron-methyl, trifloxysulfuron-sodium, and triflusulfuron-methyl. EPA is hereby reopening the comment period for 45 days.

To submit comments, or access the docket, please follow the detailed instructions provided under ADDRESSES in the Federal Register document of July 14, 2016. If you have questions,

consult the person listed under FOR FURTHER INFORMATION CONTACT.

Authority: 7 U.S.C. 136 et seq. Dated: September 14, 2016.

Yu-Ting Guilaran,

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

[FR Doc. 2016-23437 Filed 9-27-16; 8:45 am]

BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPPT-2015-0688; FRL-9947-01-OEI]

Agency Information Collection
Activities; Information Collection
Request Submitted to OMB for Review
and Approval; Comment Request;
Recordkeeping and Reporting
Requirements for Allegations of
Significant Adverse Reactions to
Human Health or the Environment

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted the following information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act: "Recordkeeping and Reporting Requirements for Allegations of Significant Adverse Reactions to Human Health or the Environment" (EPA ICR No. 1031.11, OMB Control No. 2070–0017). This is a request to renew

the approval of an existing ICR, which is currently approved through September 30, 2016. EPA did not receive any comments in response to the previously provided public review opportunity issued in the **Federal Register** of March 10, 2016 (81 FR 12730). With this submission, EPA is providing an additional 30 days for public review.

DATES: Comments must be received on or before October 28, 2016.

ADDRESSES: Submit your comments, identified by Docket ID Number EPA–HQ–OPPT–2015–0688, to both EPA and OMB as follows:

- To EPA online using http:// 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
- To 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 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:

Colby Lintner, Environmental Assistance Division (7408M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460; telephone number: (202) 554–1404; email address: *TSCA-Hotline@epa.gov*.

SUPPLEMENTARY INFORMATION:

Docket: Supporting documents, including the ICR that explains in detail the information collection activities and the related burden and cost estimates that are summarized in this document, are available in the docket for this ICR. The docket can be viewed online at http://www.regulations.gov or in person at the EPA Docket Center, West William Jefferson Clinton Bldg., Rm. 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.

ICR status: This ICR is currently scheduled to expire on September 30, 2016. Under OMB regulations, the Agency may continue to conduct or sponsor the collection of information while this submission is pending at OMB

Under PRA, 44 U.S.C. 3501 et seq., 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 are displayed either by publication in the **Federal Register** or by other appropriate means, such as on the related collection instrument or form, if applicable. The display of OMB control numbers for certain EPA regulations is consolidated in 40 CFR part 9.

Abstract: TSCA section 8(c) requires companies that manufacture, process, or distribute chemicals to maintain records of significant adverse reactions to health or the environment alleged to have been caused by such chemicals. Since section 8(c) includes no automatic reporting provision, EPA can obtain and use the information contained in company files only by inspecting those files or requiring reporting of records that relate to specific substances of concern. Therefore, under certain conditions, and using the provisions found in 40 CFR part 717, EPA may require companies to report such allegations to the Agency.

EPA uses such information on a casespecific basis to corroborate suspected adverse health or environmental effects of chemicals already under review by EPA. The information is also useful to identify trends of adverse effects across the industry that may not be apparent to any one chemical company. This ICR addresses the information reporting and recordkeeping requirements found in 40 CFR part 717.

Respondents may claim all or part of a notice as CBI. EPA will disclose

information that is covered by a CBI claim only to the extent permitted by, and in accordance with, the procedures in 40 CFR part 2.

Form Numbers: None.

Respondents/affected entities: Entities potentially affected by this ICR are companies that manufacture, process, import, or distribute in commerce chemical substances or mixtures.

Respondent's obligation to respond: Mandatory; see 40 CFR part 717. Estimated number of respondents:

13,160 (total).

Frequency of response: On occasion. Total estimated burden: 25,527 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: \$1,911,471 (per year), includes \$0 annualized capital or operation & maintenance costs.

Changes in the Estimates: There is a decrease of 1,405 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. This decrease is due to EPA's estimate of fewer potential respondents affected by the reporting requirement.

Authority: 44 U.S.C. 3501 et seq.

Courtney Kerwin,

Director, Regulatory Support Division. [FR Doc. 2016–23384 Filed 9–27–16; 8:45 am] BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-ORD-2016-0010; FRL 9953-27-OEI]

Information Collection Request Submitted to OMB for Review and Approval; Comment Request; Recordkeeping for Institutional Dual Use Research of Concern (iDURC) Policy Compliance

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted an information collection request (ICR), "Recordkeeping for Institutional Dual Use Research of Concern (iDURC) Policy Compliance" (EPA ICR No. 2530.02, OMB Control No. 2080-0082) 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 request for extension of the ICR currently approved through September 30, 2016. Public comments were previously requested via the Federal Register (81 FR 33530) on May 26, 2016 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 October 28, 2016.

ADDRESSES: Submit your comments, referencing Docket ID Number EPA—HQ—ORD—2016—0010, to (1) EPA online using www.regulations.gov (our preferred method), by email to ord.docket@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:

Brendan Doyle, Office of Research and Development, Mail Code: 8801R, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460; telephone number: 202–564–4584; email address: doyle.brendan@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: To comply with the U.S. Government Policy for Institutional Oversight of Life Sciences Dual Use Research of Concern (Policy) (www.phe.gov/s3/dualuse/Pages/default.aspx), EPA must ensure that the institutions that are subject to the Policy train their laboratory personnel and maintain records of that training. This training is specific to "dual use research of concern," and should include