

performance test guidelines to standardize the approaches to testing methods to ensure the quality and validity of the efficacy data for these types of products. The Agency attended entomology conferences, consulted with leading academics, and reviewed peer-reviewed scientific journal articles on topics related to the guideline to draft the original document. Further, EPA sought advice and recommendations from the FIFRA Scientific Advisory Panel (SAP) and the public. The SAP meeting, held on May 8–10, 2018, was announced in the **Federal Register** issue of January 26, 2018 (83 FR 3704) (FRL–9972–65). This guideline has been revised based on comments from the SAP and the public. The revisions include clarifying bait product testing, offering more flexibility in testing design, updating the replication recommendations based on statistical modeling and ease of obtaining pests, and refining the statistical analyses recommendations. The Agency is also making available in the docket a Response to Comments document that addresses issue raised in the public comment submission.

Authority: 7 U.S.C. 136 *et seq.*; 15 U.S.C. 2601 *et seq.*; 21 U.S.C. 301 *et seq.*

Dated: September 24, 2019.

Alexandra Dapolito Dunn,

Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OA–2019–0370; FRL–9047–2]

Proposed Information Collection Request; Comment Request; Final Rule at 40 CFR Part 8: Environmental Impact Assessment of Nongovernmental Activities in Antarctica (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency is planning to submit an information collection request (ICR), “Final Rule at 40 CFR part 8: Environmental Impact Assessment of Nongovernmental Activities in Antarctica” (EPA ICR No. 1808.09, OMB Control No. 2020–0007) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act. Before doing so, the 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 May 31, 2020. 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 Friday, November 29, 2019.

ADDRESSES: Submit your comments, referencing Docket ID No. EPA–HQ–OA–2019–0370 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.

The 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 or other information whose disclosure is restricted by statute.

FOR FURTHER INFORMATION CONTACT: Julie Roemele, NEPA Compliance Division, Office of Federal Activities, Mail Code 2203A, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: 202–564–5632; fax number: 202–564–0070; email address: roemele.julie@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, the 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. The 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, the 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: The EPA’s regulations at 40 CFR part 8, Environmental Impact Assessment of Nongovernmental Activities in Antarctica (Rule), were promulgated pursuant to the Antarctic Science, Tourism, and Conservation Act of 1996 (Act), 16 U.S.C. 2401 *et seq.*, as amended, 16 U.S.C. 2403a, which implements the Protocol on Environmental Protection (Protocol) to the Antarctic Treaty of 1959 (Treaty). The Rule provides for assessment of the environmental impacts of nongovernmental activities in Antarctica, including tourism, for which the United States is required to give advance notice under Paragraph 5 of Article VII of the Treaty, and for coordination of the review of information regarding environmental impact assessments received from other Parties under the Protocol. The requirements of the Rule apply to operators of nongovernmental expeditions organized or proceeding from the territory of the United States to Antarctica and include commercial and non-commercial expeditions. Expeditions may include ship-based tours; yacht, skiing or mountaineering expeditions; privately funded research expeditions; and other nongovernmental activities. The Rule does not apply to individual U.S. citizens or groups of citizens planning travel to Antarctica on an expedition for which they are not acting as an operator. (Operators, for example, typically acquire use of vessels or aircraft, hire expedition staff, plan itineraries, and undertake other organizational responsibilities.) The rule provides nongovernmental operators with the specific requirements they need to meet in order to comply with the requirements of Article 8 and Annex I to the Protocol. The provisions of the Rule are intended to ensure that potential environmental effects of nongovernmental activities undertaken in Antarctica are appropriately identified and considered by the operator during the planning process and that to the extent practicable

appropriate environmental safeguards which would mitigate or prevent adverse impacts on the Antarctic environment are identified by the operator.

Environmental Documentation.

Persons subject to the Rule must prepare environmental documentation to support the operator's determination regarding the level of environmental impact of the proposed expedition. Environmental documentation includes a Preliminary Environmental Review Memorandum (PERM), an Initial Environmental Evaluation (IEE), or a Comprehensive Environmental Evaluation (CEE). The environmental document is submitted to the Office of Federal Activities (OFA). If the operator determines that an expedition may have: (1) Less than a minor or transitory impact, a PERM needs to be submitted no later than 180 days before the proposed departure to Antarctica; (2) no more than minor or transitory impacts, an IEE needs to be submitted no later than 90 days before the proposed departure; or (3) more than minor or transitory impacts, a CEE needs to be submitted. Operators who anticipate such activities are encouraged to consult with EPA as soon as possible regarding the date for submittal of the CEE. (Article 3(4), of Annex I of the Protocol requires that draft CEEs be distributed to all Parties and the Committee for Environmental Protection 120 days in advance of the next Antarctic Treaty Consultative Meeting at which the CEE may be addressed.)

The Protocol and the Rule also require an operator to employ procedures to assess and provide a regular and verifiable record of the actual impacts of an activity which proceeds on the basis of an IEE or CEE. The record developed through these measures needs to be designed to: (a) Enable assessments to be made of the extent to which environmental impacts of nongovernmental expeditions are consistent with the Protocol; and (b) provide information useful for minimizing and mitigating those impacts and, where appropriate, on the need for suspension, cancellation, or modification of the activity. Moreover, an operator needs to monitor key environmental indicators for an activity proceeding based on a CEE. An operator may also need to carry out monitoring in order to assess and verify the impact of an activity for which an IEE would be prepared. For activities that require an IEE, an operator should be able to use procedures currently being voluntarily utilized by operators to provide the required information. Should an activity require a CEE, the operator should

consult with the EPA to: (a) Identify the monitoring regime appropriate to that activity, and (b) determine whether and how the operator might utilize relevant monitoring data collected by the U.S. Antarctic Program. OFA would consult with the National Science Foundation and other interested Federal agencies regarding the monitoring regime.

In cases of emergency related to the safety of human life or of ships, aircraft, equipment and facilities of high value, or the protection of the environment which would require an activity to be undertaken without completion of the documentation procedures set out in the Rule, the operator would need to notify the Department of State within 15 days of any activities which would have otherwise required preparation of a CEE, and provide a full explanation of the activities carried out within 45 days of those activities. (During the time the Interim Final and Final Rules have been in effect, there were no emergencies requiring notification by U.S. operators. An Interim Final Rule was in effect from April 30, 1997, until replaced on December 6, 2001, by the Final Rule).

Environmental documents (e.g., PERM, IEE, CEE) are submitted to OFA. Environmental documents are reviewed by OFA, in consultation with the National Science Foundation and other interested Federal agencies and made available to other Parties and the public as required under the Protocol or otherwise requested. OFA notifies the public of document availability via the World Wide Web at: <https://www.epa.gov/international-cooperation/receipt-environmental-impact-assessments-eias-regarding-nongovernmental>.

The types of nongovernmental activities currently being carried out (e.g., ship-based tours, land-based tours, flights, and privately funded research expeditions) are typically unlikely to have impacts that are more than minor or transitory, thus an IEE is the typical level of environmental documentation submitted. For the 1997–1998 through 2018–2019 austral summer seasons during the time the Rule has been in effect, all respondents submitted IEEs except for three PERMs. Paperwork reduction provisions in the Rule that are used by the operators include: (a) Incorporation of material in the environmental document by referring to it in the IEE, (b) inclusion of all proposed expeditions by one operator within one IEE; (c) use of one IEE to address expeditions being carried out by more than one operator; and (d) use of multi-year environmental documentation to address proposed

expeditions for a period of up to five consecutive austral summer seasons.

Coordination of Review of Information Received From Other Parties to the Treaty. The Rule also provides for the coordination of review of information received from other Parties and the public availability of that information including: (1) A description of national procedures for considering the environmental impacts of proposed activities; (2) an annual list of any IEEs and any decisions taken in consequence thereof; (3) significant information obtained and any action taken in consequence thereof with regard to monitoring from IEEs to CEEs; and (4) information in a final CEE. This provision fulfills the United States' obligation to meet the requirements of Article 6 of Annex I to the Protocol. The Department of State is responsible for coordination of these reviews of drafts with interested Federal agencies, and for public availability of documents and information. This portion of the Rule does not impose paperwork requirements on any nongovernmental person subject to U.S. regulation.

Form Numbers: None.

Respondents/affected entities: Entities potentially affected by this action are all nongovernmental operators with activities in Antarctica, including tour operators, for which the United States is required to give advance notice under paragraph 5 of Article VII of the Antarctic Treaty of 1959; this includes all nongovernmental expeditions to and within Antarctica organized in or proceeding from the territory of the United States.

Respondent's obligation to respond: Mandatory (40 CFR part 8).

Estimated number of respondents: 25 (total).

Frequency of response: Annual.

Total estimated burden: 1,544 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: \$133,780 (per year), includes \$0 annualized capital or operation & maintenance costs.

Changes in Estimates: There is an increase of 330 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. This increase is the result of a change to the number of operators that the EPA anticipates will submit environmental documentation as well as the inclusion of a potential PERM, CEE and Emergency Report submitted by every three years.

Dated: September 24, 2019.

Robert Tomiak,

Director, Office of Federal Activities.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0720; FRL–9997–21]

Registration Review; Draft Human Health and/or Ecological Risk Assessments for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA’s draft human health and/or ecological risk assessments for the registration review of alkylbenzene sulfonates, cyproconazole, etoxazole, fenamidone, fenbutatin-oxide, fluzifop-p-butyl, flumetralin, MCPB and salts, mecoprop (MCP-p), oxyfluorfen, pinoxaden, pyraclostrobin, and pyraflufen-ethyl.

DATES: Comments must be received on or before November 29, 2019.

ADDRESSES: Submit your comments, to the docket identification (ID) number for the specific pesticide of interest provided in the Table in Unit IV., by one of the following methods:

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

Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute.

- *Mail:* OPP Docket, Environmental Protection Agency Docket Center (EPA/DC), (28221T), 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001.

- *Hand Delivery:* To make special arrangements for hand delivery or delivery of boxed information, please follow the instructions at <http://www.epa.gov/dockets/contacts.html>.

Additional instructions on commenting or visiting the docket, along with more information about dockets generally, are available at <http://www.epa.gov/dockets>.

FOR FURTHER INFORMATION CONTACT: For pesticide specific information contact: The Chemical Review Manager for the pesticide of interest identified in the Table in Unit IV.

For general questions on the registration review program, contact: Melanie Biscoe, Pesticide Re-Evaluation Division (7508P), Office of Pesticide

Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (703) 305–7106; email address: biscoe.melanie@epa.gov.

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, farm worker, 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 Chemical Review Manager identified in the Table in Unit IV.

B. What should I consider as I prepare my comments for EPA?

1. *Submitting CBI.* Do not submit this information to EPA through [regulations.gov](http://www.regulations.gov) or email. Clearly mark the part or all of the information that you claim to be CBI. For CBI information in a disk or CD–ROM that you mail to EPA, mark the outside of the disk or CD–ROM as CBI and then identify electronically within the disk or CD–ROM the specific information that is claimed as CBI. In addition to one complete version of the comment that includes information claimed as CBI, a copy of the comment that does not contain the information claimed as CBI must be submitted for inclusion in the public docket. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

2. *Tips for preparing your comments.* When preparing and submitting your comments, see the commenting tips at <http://www.epa.gov/dockets/comments.html>.

3. *Environmental justice.* EPA seeks to achieve environmental justice, the fair treatment and meaningful involvement of any group, including minority and/or low income populations, in the development, implementation, and enforcement of environmental laws, regulations, and policies. To help address potential environmental justice issues, the Agency seeks information on any groups or segments of the population who, as a result of their location, cultural practices, or other factors, may have atypical or disproportionately high and adverse

human health impacts or environmental effects from exposure to the pesticides discussed in this document, compared to the general population.

II. Background

Registration review is EPA’s periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. As part of the registration review process, the Agency has completed comprehensive draft human health and/or ecological risk assessments for all pesticides listed in the Table in Unit IV. After reviewing comments received during the public comment period, EPA may issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments and may request public input on risk mitigation before completing a proposed registration review decision for the pesticides listed in the Table in Unit IV. Through this program, EPA is ensuring that each pesticide’s registration is based on current scientific and other knowledge, including its effects on human health and the environment.

III. Authority

EPA is conducting its registration review of the chemicals listed in the Table in Unit IV pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. FIFRA section 3(g) provides, among other things, that the registrations of pesticides are to be reviewed every 15 years. Under FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5) (7 U.S.C. 136a(c)(5)). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

IV. What action is the Agency taking?

Pursuant to 40 CFR 155.58, this notice announces the availability of EPA’s human health and/or ecological risk assessments for the pesticides shown in the following table and opens a 60-day public comment period on the risk assessments.