compliance may only be achieved by submission of the data/information described in Table 2 of Unit II.

VI. Status of Products That Become Suspended

Your product will remain suspended, however, until the Agency determines you are in compliance with the requirements which are the bases of this notice and so informs you in writing.

After the suspension becomes final and effective, the registrants subject to this notice, including all supplemental registrants of products listed in Table 1 of Unit II., may not legally distribute, sell, use, offer for sale, hold for sale, ship, deliver for shipment, or receive and (having so received) deliver or offer to deliver, to any person, the products listed in Table 1 of Unit II. Persons other than the registrants subject to this notice, as defined in the preceding sentence, may continue to distribute, sell, use, offer for sale, hold for sale, ship, deliver for shipment, or receive and (having so received) deliver or offer to deliver, to any person, the products listed in Table 1 of Unit II. Nothing in this notice authorizes any person to distribute, sell, use, offer for sale, hold for sale, ship, deliver for shipment, or receive and (having so received) deliver or offer to deliver, to any person, the products listed in Table 1 of Unit II. in any manner which would have been unlawful prior to the suspension.

If the registrations for your products, listed in Table 1 of Unit II., are currently suspended as a result of failure to comply with another FIFRA section 3(c)(2)(B) DCI notice or FIFRA Section 4 Data Requirements notice, this notice, when it becomes a final and effective order of suspension, will be in addition to any existing suspension, *i.e.,* all requirements which are the bases of the suspension must be satisfied before the registration will be reinstated.

It is the responsibility of the basic registrant to notify all supplementary registered distributors of a basic registered product that this suspension action also applies to their supplementary registered products. The basic registrant may be held liable for violations committed by their distributors.

Any questions about the requirements and procedures set forth in this notice or in the subject FIFRA section 3(c)(2)(B) DCI notice, should be addressed to the person listed under **FOR FURTHER INFORMATION CONTACT**.

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

Dated: August 19, 2020.

Anita Pease,

Director, Antimicrobials Division, Office of Pesticide Programs.

[FR Doc. 2020–19370 Filed 9–1–20; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0750; FRL–10012–60]

Pesticide Registration Review; Proposed Interim Decisions for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's proposed interim registration review decisions and opens a 60-day public comment period on the proposed interim decisions for the following pesticides: 1, 4-Dimethylnapthalene and 2,6-Diisopropylnaphthalene, acequinocyl, *Bacillus cereus* strain BP01, cypermethrins, dithiopyr, etridiazole, fenamidone, fenbutatin-oxide, fenpropimorph, fenpyroximate, flonicamid, flumetralin, flumioxazin, hypochlorous acid, inorganic halides, MCPB, *Metarhizium anisopliae,* metolachlor/S-metolachlor, *Pantoea agglomerans* strain C9–1, *Pantoea agglomerans* strain E325, propanil, terbacil, triclopyr.

DATES: Comments must be received on or before November 2, 2020.

ADDRESSES: Submit your comments, identified by 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*.

Please note that due to the public health emergency the EPA Docket Center (EPA/DC) and Reading Room was closed to public visitors on March 31, 2020. Our EPA/DC staff will continue to provide customer service via email, phone, and webform. For further information on EPA/DC services, docket contact information and the current status of the EPA/DC and Reading Room, please visit *https://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 information 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 for the pesticide of interest 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* or email. Clearly mark the part or all of the information that you claim to be CBI. For CBI information on 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.

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 proposed interim decisions for all 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. Section 3(g) of FIFRA 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 proposed interim registration review decisions for the pesticides shown in the table below and opens a 60-day public comment period on the proposed interim registration review decisions.

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
1, 4-Dimethylnapthalene and 2,6-Diisopropylnaphthalene, Case Number 6029.	EPA–HQ–OPP–2012–0670	Joseph Mabon, *mabon.joseph@epa.gov*, (703) 347–0177.
Acequinocyl, Case Number 7621	EPA–HQ–OPP–2015–0203	Sergio Santiago, *santiago.sergio@epa.gov*, (703) 347–8606.
Bacillus cereus strain BP01, Case Number 6053	EPA–HQ–OPP–2011–0493	Alexandra Boukedes, *boukedes.alexandra@epa.gov*, (703) 347–0305.
Cypermethrins, Case Number 2130	EPA–HQ–OPP–2012–0167	Susan Bartow, *bartow.susan@epa.gov*, (703) 603–0065.
Dithiopyr, Case Number 7225	EPA–HQ–OPP–2013–0750	Veronica Dutch, *dutch.veronica@epa.gov*, (703) 308–8585.
Etridiazole, Case Number 0009	EPA–HQ–OPP–2014–0414	Jonathan Williams, *williams.jonathanr@epa.gov*, (703) 347–0670.
Fenamidone, Case 7033	EPA–HQ–OPP–2014–0048	Christian Bongard, *bongard.christian@epa.gov*, (703) 347–0337.
Fenbutatin Oxide, Case Number 0245	EPA–HQ–OPP–2009–0841	Lauren Weissenborn, *weissenborn.lauren@epa.gov*, (703) 347–8601.
Fenpropimorph, Case Number 5112	EPA–HQ–OPP–2014–0404	Peter Bergquist, *bergquist.peter@epa.gov*, (703) 347–8563.
Fenpyroximate, Case Number 7432	EPA–HQ–OPP–2014–0572	Carolyn Smith, *smith.carolyn@epa.gov*, (703) 347–8325.
Flonicamid, Case Number 7436	EPA–HQ–OPP–2014–0777	Alexandra Feitel, *feitel.alexandra@epa.gov*, (703) 347–8631.
Flumetralin, Case Number 4119	EPA–HQ–OPP–2015–0076	Theodore Varns, *varns.theodore@epa.gov*, (703) 347–8589.
Flumioxazin, Case Number 7244	EPA–HQ–OPP–2011–0176	Susan Bartow, *bartow.susan@epa.gov*, (703) 603–0065.
Hypochlorous Acid, Case Number 5090	EPA–HQ–OPP–2020–0244	Jessie Bailey, *bailey.jessica@epa.gov*, (703) 347–0148.
Inorganic Halides, Case Number 4051	EPA–HQ–OPP–2009–0168	Erin Dandridge, *dandridge.erin@epa.gov*, (703) 347–0185.
MCPB, Case Number 2365	EPA–HQ–OPP–2014–0181	Steven R. Peterson, *peterson.stevenr@epa.gov*, (703) 347–0755.
Metarhizium anisopliae, Case Number 6024	EPA–HQ–OPP–2009–0510	Susanne Cerrelli, *cerrelli.susanne@epa.gov*, (703) 308–8077.
Metolachlor/S-metolachlor, Case Number 0001	EPA–HQ–OPP–2014–0772	Ana Pinto, *pinto.ana@epa.gov*, (703) 347–8421.
Pantoea agglomerans strain C9–1, Case Number 6506.	EPA–HQ–OPP–2017–0172	Bibiana Oe, *oe.bibiana@epa.gov*, (703) 347–8162.
Pantoea agglomerans strain E325, Case Number 6507.	EPA–HQ–OPP–2017–0172	Bibiana Oe, *oe.bibiana@epa.gov*, (703) 347–8162.
Propanil, Case Number 0226	EPA–HQ–OPP–2015–0052	Tiffany Green, *green.tiffany@epa.gov*, (703) 347–0314.
Terbacil, Case Number 0039	EPA–HQ–OPP–2011–0054	Alexandra Feitel, *feitel.alexandra@epa.gov*, (703) 347–8631.
Triclopyr, Case Number 2710	EPA–HQ–OPP–2014–0576	Andy Muench, *muench.andrew@epa.gov*, (703) 347–8263.

The registration review docket for a pesticide includes earlier documents related to the registration review case. For example, the review opened with a Preliminary Work Plan, for public comment. A Final Work Plan was placed in the docket following public comment on the Preliminary Work Plan.

The documents in the dockets describe EPA's rationales for conducting additional risk assessments for the registration review of the pesticides included in the tables in Unit IV, as well as the Agency's subsequent risk findings and consideration of possible risk mitigation measures. These proposed interim registration review decisions are supported by the rationales included in those documents. Following public comment, the Agency will issue interim or final registration review decisions for the pesticides listed in Table 1 in Unit IV.

The registration review final rule at 40 CFR 155.58(a) provides for a minimum 60-day public comment period on all proposed interim registration review decisions. This comment period is intended to provide an opportunity for public input and a mechanism for initiating any necessary amendments to the proposed interim decision. All comments should be submitted using the methods in **ADDRESSES** and must be received by EPA on or before the closing date. These comments will become part of the docket for the pesticides included in the Tables in Unit IV. Comments received after the close of the comment period will be marked ''late.'' EPA is not required to consider these late comments.

The Agency will carefully consider all comments received by the closing date and may provide a ''Response to Comments Memorandum'' in the docket. The interim registration review decision will explain the effect that any

comments had on the interim decision and provide the Agency's response to significant comments.

Background on the registration review program is provided at: *http://www.epa.gov/pesticide-reevaluation.*

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

Dated: July 23, 2020.

Mary Reaves,

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

[FR Doc. 2020–19374 Filed 9–1–20; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OW–2020–0433; FRL–10014–43–OW]

Proposed Information Collection Request; Comment Request; Public Notification Requirements for Combined Sewer Overflows in the Great Lakes Basin (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) is planning to submit an information collection request (ICR), ''Public Notification Requirements for Combined Sewer Overflows in the Great Lakes Basin (Renewal)'' (EPA ICR No. 2562.02, OMB Control No. 2040–0293) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act (PRA). Before doing so, EPA is soliciting public comments on specific aspects of the proposed information collection. This is a proposed extension of the ICR which is currently approved through April 30, 2021. 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 November 2, 2020.

ADDRESSES: You may send comments, identified by Docket ID No. EPA–HQ–OW–2020–0433, by any of the following methods:

- *Federal eRulemaking Portal: https://www.regulations.gov/* (our preferred method). Follow the online instructions for submitting comments.
- *Email: Baehr.Joshua@epa.gov.* Include Docket ID No. EPA–HQ–OW–2020–0433 in the subject line of the message.

Instructions: All submissions received must include the Docket ID No. for this rulemaking. Comments received may be posted without change to *https://www.regulations.gov/,* including any personal information provided. For detailed instructions on sending comments and additional information on the rulemaking process, see the ''How do I submit written comments?'' heading of the **SUPPLEMENTARY INFORMATION** section of this document. Out of an abundance of caution for members of the public and our staff, the EPA Docket Center and Reading Room are closed to the public, with limited exceptions, to reduce the risk of transmitting COVID–19. Our Docket Center staff will continue to provide remote customer service via email, phone, and webform. We encourage the public to submit comments via *https://www.regulations.gov* or email, as there may be a temporary delay in processing mail and faxes. Hand deliveries may be received by scheduled appointment only. For further information on EPA Docket Center services and the current status, please visit us online at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Joshua Baehr, National Program Branch, Water Permits Division, OWM Mail Code: 4203M, Environmental Protection Agency, 1201 Constitution Ave. NW, Washington, DC 20460; telephone number: (202) 564–2277; email address: *Baehr.Joshua@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. How do I submit written comments?

Submit your comments, identified by Docket ID No. EPA–HQ–OW–2020–0433, at *https://www.regulations.gov* (our preferred method), or the other methods identified in the **ADDRESSES** section. Once submitted, comments cannot be edited or removed from the docket. The EPA may publish any comment received to its public docket. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by a written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. The EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.* on the web, cloud, or other file sharing system). For additional submission methods, the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit *https://www.epa.gov/dockets/commenting-epa-dockets.*

The EPA is temporarily suspending its Docket Center and Reading Room for public visitors to reduce the risk of transmitting COVID–19. Written comments submitted by mail are temporarily suspended and no hand deliveries will be accepted. Our Docket Center staff will continue to provide remote customer service via email, phone, and webform. We encourage the public to submit comments via *https://www.regulations.gov.* For further information and updates on EPA Docket Center services, please visit us online at *https://www.epa.gov/dockets.*

The EPA continues to carefully and continuously monitor information from the Centers for Disease Control and Prevention (CDC), local area health departments, and our Federal partners so that we can respond rapidly as conditions change regarding COVID–19.

II. Executive Summary

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 *https://www.regulations.gov.* The telephone number for the Docket Center is 202–566–1744. For additional information about EPA's public docket, visit *https://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; (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: This ICR calculates the incremental increase in burden and