TABLE—DRAFT RISK ASSESSMENTS BEING MADE AVAILABLE FOR PUBLIC COMMENT

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Carboxin/Oxycarboxin, Case 0012	EPA–HQ–OPP–2004–0124	Tiffany Green, *green.tiffany@epa.gov*, (703) 347–0314.
Famoxadone, Case 7038	EPA–HQ–OPP–2015–0094	Christina Scheltema, *scheltema.christina@epa.gov*, (703) 305–8401.
Fenpropimorph, Case 5112	EPA–HQ–OPP–2014–0404	Peter Bergquist, *bergquist.peter@epa.gov*, (703) 347–8563.
Irgarol, Case 5031	EPA–HQ–OPP–2010–0003	SanYvette Williams, *williams.sanyvette@epa.gov*, (703) 305–7702.
Naphthalene Acetic Acid and its Salts, Ester, and Acetamide (NAA), Case 0379.	EPA–HQ–OPP–2014–0773	Linsey Walsh, *walsh.linsey@epa.gov*, (703) 347–0588.
Propargite, Case 0243	EPA–HQ–OPP–2014–0131	Wilhelmena Livingston, *livingston.wilhelmena@epa.gov*, (703) 308–8025.
Telone (1,3-D), Case 0328	EPA–HQ–OPP–2013–0154	Michelle Nolan, *nolan.michelle@epa.gov*, (703) 347–0258.
Triallate, Case 2695	EPA–HQ–OPP–2014–0573	Katherine St. Clair, *stclair.Katherine@epa.gov*, (703) 347–8778.
Triticonazole, Case 7036	EPA–HQ–OPP–2015–0602	Christian Bongard, *bongard.christian@epa.gov*, (703) 347–0337.

Pursuant to 40 CFR 155.53(c), EPA is providing an opportunity, through this notice of availability, for interested parties to provide comments and input concerning the Agency's draft human health and/or ecological risk assessments for the pesticides listed in the Table in Unit IV. The Agency will consider all comments received during the public comment period and make changes, as appropriate, to a draft human health and/or ecological risk assessment. EPA may then issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments.

Information submission requirements. Anyone may submit data or information in response to this document. To be considered during a pesticide's registration review, the submitted data or information must meet the following requirements:

- To ensure that EPA will consider data or information submitted, interested persons must submit the data or information during the comment period. The Agency may, at its discretion, consider data or information submitted at a later date.
- The data or information submitted must be presented in a legible and useable form. For example, an English translation must accompany any material that is not in English and a written transcript must accompany any information submitted as an audiographic or videographic record. Written material may be submitted in paper or electronic form.
- Submitters must clearly identify the source of any submitted data or information.
- Submitters may request the Agency to reconsider data or information that the Agency rejected in a previous review. However, submitters must explain why they believe the Agency should reconsider the data or information in the pesticide's registration review.

As provided in 40 CFR 155.58, the registration review docket for each pesticide case will remain publicly accessible through the duration of the registration review process; that is, until all actions required in the final decision on the registration review case have been completed.

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

Dated: January 27, 2020.

Mary Reaves,

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

[FR Doc. 2020–02041 Filed 2–3–20; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0720; FRL–10004–43]

Pesticide Registration Review; Pesticide Dockets Opened for Review and Comment; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of the EPA's preliminary work plans for the following chemicals: Methoprene, pyridalyl, and spirotetramat. With this document, the EPA is opening the public comment period for registration review for these chemicals.

DATES: Comments must be received on or before April 6, 2020.

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 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 the EPA?

1. *Submitting CBI.* Do not submit this information to the EPA through *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 the 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 the 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. Registration review dockets contain information that will assist the public in understanding the types of information and issues that the agency may consider during the course of registration reviews. As part of the registration review process, the Agency has completed preliminary workplans for all pesticides listed in the Table in Unit IV. Through this program, the 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

The 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. Registration Reviews

A. What action is the Agency taking?

A pesticide's registration review begins when the agency establishes a docket for the pesticide's registration review case and opens the docket for public review and comment. Pursuant to 40 CFR 155.50, this notice announces the availability of the EPA's preliminary work plans for the pesticides shown in the following table and opens a 60-day public comment period on the work plans.

TABLE—REGISTRATION REVIEW CASES

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Methoprene, Kinoprene, and Hydroprene, Case No. 0030.	EPA–HQ–OPP–2013–0586	Cody Kendrick, *kendrick.cody@epa.gov,* (703) 347–0468.
Pyridalyl, Case No. 7451	EPA–HQ–OPP–2019–0378	Sergio Santiago, *santiago.sergio@epa.gov,* (703) 347–8606.
Spirotetramat, Case No. 7452	EPA–HQ–OPP–2019–0033	Darius Stanton, *stanton.darius@epa.gov,* (703) 347–0433.

B. Docket Content

The registration review docket contains information that the agency may consider in the course of the registration review. The agency may include information from its files including, but not limited to, the following information:

- An overview of the registration review case status.
- A list of current product registrations and registrants.
- **Federal Register** notices regarding any pending registration actions.
- **Federal Register** notices regarding current or pending tolerances.
- Risk assessments.
- Bibliographies concerning current registrations.
- Summaries of incident data.
- Any other pertinent data or information.

Each docket contains a document summarizing what the agency currently knows about the pesticide case and a preliminary work plan for anticipated data and assessment needs. Additional documents provide more detailed information. During this public comment period, the agency is asking that interested persons identify any additional information they believe the agency should consider during the registration reviews of these pesticides. The agency identifies in each docket the areas where public comment is specifically requested, though comment in any area is welcome.

The registration review final rule at 40 CFR 155.50(b) provides for a minimum 60-day public comment period on all preliminary registration review work plans. This comment period is intended to provide an opportunity for public input and a mechanism for initiating any necessary changes to a pesticide's workplan. All comments should be submitted using the methods in **ADDRESSES** and must be received by the EPA on or before the closing date. These comments will become part of the docket for the pesticides included in the Table in Unit IV. Comments received after the close of the comment period will be marked ''late.'' The 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 final registration review work plan will explain the effect that any comments had on the final work plan 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: January 27, 2020.

Mary Reaves,

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

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