TABLE—REGISTRATION REVIEW INTERIM DECISIONS BEING ISSUED—Continued

Registration review case name and number	Docket ID No.	Chemical review manager and contact information
Diquat dibromide, Case Number 0288	EPA–HQ–OPP–2009–0846	Jordan Page, *page.jordan@epa.gov* (703) 347–0467.
Fluthiacet-methyl, Case Number 7280	EPA–HQ–OPP–2013–0285	Eric Fox, *fox.ericm@epa.gov* (703) 347–0104.
Hydramethylnon, Case Number 2585	EPA–HQ–OPP–2012–0869	Carolyn Smith, *smith.carolyn@epa.gov* (703) 347–8325.
Inorganic polysulfides (also known as calcium polysulfide or lime sulfur) Case Number 4054.	EPA–HQ–OPP–2016–0102	Katherine St. Clair, *stclair.katherine@epa.gov* (703) 347–8778.
IR3535, Case Number 6046	EPA–HQ–OPP–2014–0106	Alexandra Boukedes, *boukedes.alexandra@epa.gov* (703) 347–0305.
Lufenuron, Case Number 7627	EPA–HQ–OPP–2015–0098	Andy Muench, *muench.andrew@epa.gov* (703) 347–8263.
o-Benzyl-p-Chlorophenol (OBPCP), Case Number 2045.	EPA–HQ–OPP–2011–0423	Erin Dandridge, *dandridge.erin@epa.gov* (703) 347–0185.
Octenol, Case Number 6033	EPA–HQ–OPP–2012–0940	Joseph Mabon, *mabon.joseph@epa.gov* (703) 347–0177.
p-Methane-3,8-diol (PMD), Case Number 6017	EPA–HQ–OPP–2015–0693	Joseph Mabon, *mabon.joseph@epa.gov* (703) 347–0177.
Potato Leaf Roll Virus Resistance Gene, Case Number 6505.	EPA–HQ–OPP–2012–0416	Michael Glikes, *glikes.michael@epa.gov* (703) 305–6231.
Starlicide (DRC–1339), Case Number 2610	EPA–HQ–OPP–2011–0696	Nathan Sell, *sell.nathan@epa.gov* (703) 347–8020.
Trifluralin, Case Number 0179	EPA–HQ–OPP–2012–0417	Matthew Khan, *khan.matthew@epa.gov* (703) 347–8613.
Tri-n Butyl Tetradecyl Phosphonium Chloride (TTPC), Case Number 5111.	EPA–HQ–OPP–2011–0952	Daniel Halpert, *halpert.daniel@epa.gov* (703) 347–0133.
Uniconazole-P, Case Number 7007	EPA–HQ–OPP–2015–0729	Jaclyn Pyne, *pyne.jaclyn@epa.gov* (703) 347–0445.
Zinc and Zinc Salts, Case Number 4099	EPA–HQ–OPP–2009–0011	Michael McCarroll, *mccarroll.michael@epa.gov* (703) 347–0147.
Zoxamide, Case Number 7032	EPA–HQ–OPP–2014–0391	Sergio Santiago, *santiago.sergio@epa.gov* (703) 347–8606.

The proposed interim registration review decisions for the chemicals in the table above were posted to the docket and the public was invited to submit any comments or new information. EPA addressed the comments or information received during the 60-day comment period for the proposed interim decisions in the discussion for each pesticide listed in the table. Comments from the 60-day comment period that were received may or may not have affected the Agency's interim decision. Pursuant to 40 CFR 155.58(c), the registration review case docket for the chemicals listed in the Table will remain open until all actions required in the interim decision have been completed.

This document also announces the closure of the registration review case for 1-(3-chloroallyl)-3,5,7-triaza-1-azoniaadamantane (CTAC) (Case Number 3069, Docket ID Number EPA–HQ–OPP–2010–0004) because the last U.S. registrations for these pesticides have been canceled.

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.

[FR Doc. 2020–02045 Filed 1–31–20; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0750; FRL–10004–38]

Pesticide Registration Review; Proposed Interim Decisions for Several Neonicotinoid 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 acetamiprid, clothianidin, dinotefuran, imidacloprid, and thiamethoxam.

DATES: Comments must be received on or before April 3, 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.*

Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is 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 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 Table 1, and opens a 60-day public comment period on the proposed interim registration review decisions.

TABLE 1—PROPOSED INTERIM DECISIONS

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Acetamiprid, Case Number 7617	EPA–HQ–OPP–2012–0329	Jonathan Williams, *Williams.jonathanr@epa.gov,* 703–347–0670.
Clothianidin, Case Number 7620	EPA–HQ–OPP–2011–0865	Matthew Khan, *Khan.matthew@epa.gov,* 703–347–8613.
Dinotefuran, Case Number 7441	EPA–HQ–OPP–2011–0920	Steven Snyderman, *Snyderman.steven@epa.gov,* 703–347–0249.
Imidacloprid, Case Number 7605 ...	EPA–HQ–OPP–2008–0844	Steven Snyderman, *Snyderman.steven@epa.gov,* 703–347–0249.
Thiamethoxam, Case Number 7641	EPA–HQ–OPP–2011–0581	Matthew Khan, *Khan.matthew@epa.gov,* 703–347–8613.

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: January 22, 2020.

Mary Reaves,

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

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