



BILLING CODE 6560-50-P

## ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2017-0750; FRL-10001-70]

### **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: 2-phenethyl propionate, ammonium bromide, azadirachtin, *Bacillus sphaericus*, chloropicrin, *Colletotrichum gloeosporioides*, Cuelure, cyazofamid, dazomet, Extract of Reynoutria sachalinensis, fluroxypyr, glycolic acid and salts, gonadotropin releasing hormone (GnRH), Harpin Proteins, iodine and iodophors, metam sodium and metam potassium, methyl isothiocyanate (MITC), pelargonic acid salts and esters, phenmedipham, *Pythium oligandrum* DV 74, sethoxydim, tetraacetythylenediamine (TAED), thymol, tralopyril, and triclosan.

**DATES:** Comments must be received on or before **[INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE *FEDERAL REGISTER*]**.

**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](mailto: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.

This notice also announces the availability of EPA’s human health and/or ecological risk assessments for the pesticides cyazofamid, ammonium bromide, glycolic acid and salts, pelargonic acid salts and esters, sethoxydim, TAED, and thymol and opens a 60-day public comment period on the risk assessments. Additionally, this notice announces the availability of EPA’s preliminary workplan and human health and/or ecological risk assessments for GnRH.

**Table 1. Proposed Interim Decisions**

<b>Registration Review Case Name and Number</b>	<b>Docket ID Number</b>	<b>Chemical Review Manager and Contact Information</b>
2-Phenethyl Propionate Case 3110	EPA-HQ-OPP-2010-0714	Cody Kendrick <i>kendrick.cody@epa.gov</i> (703) 347-0468
Ammonium bromide Case 5002	EPA-HQ-OPP-2012-0683	Stephen Savage <i>savage.stephen@epa.gov</i> (703) 347-0345
Azadirachtin Case 6021	EPA-HQ-OPP-2008-0632	Joseph Mabon <i>mabon.joseph@epa.gov</i> (703) 347-0177
Bacillus sphaericus	EPA-HQ-OPP-2013-0116	Alexandra Boukedes

Case 6052		<i>boukedes.alexandra@epa.gov</i> (703) 347-0305
Chloropicrin Case 0040	EPA-HQ-OPP-2013-0153	Samantha Thomas <i>thomas.samantha@epa.gov</i> (703) 347-0514
Colletotrichum gloeosporioides Case 4103	EPA-HQ-OPP-2016-0685	Joseph Mabon <i>mabon.joseph@epa.gov</i> (703) 347-0177
Cuelure [2-butanone, 4-(4- (acetyloxy)phenyl)-, acetate] Case 6201	EPA-HQ-OPP-2017-0221	Bibiana Oe <i>oe.bibiana@epa.gov</i> (703) 347-8162
Cyazofamid Case 7056	EPA-HQ-OPP-2015-0128	Tiffany Green <i>green.tiffany@epa.gov</i> (703) 347-0341
Dazomet Case 2135	EPA-HQ-OPP-2013-0080	Katherine St. Clair <i>stclair.katherine@epa.gov</i> (703) 347-8778
Extract of Reynoutria sachalinensis Case 6030	EPA-HQ-OPP-2016-0232	Alexandra Boukedes <i>boukedes.alexandra@epa.gov</i> (703) 347-0305
Fluroxypyr Case 7248	EPA-HQ-OPP-2014-0570	Eric Fox <i>fox.ericm@epa.gov</i> (703) 347-0104
Glycolic Acid and Salts Case 4045	EPA-HQ-OPP-2011-0422	Michael McCarroll <i>mccarroll.michael@epa.gov</i> (703) 347-0147
Gonadotropin Releasing Hormone (GnRH) Case 7800	EPA-HQ-OPP-2018-0798	Jaclyn Pyne <i>pyne.jaclyn@epa.gov</i> (703) 347-0445
Harpin Protein and Harpin $\alpha\beta$ Protein (Harpin Proteins) Case 6010	EPA-HQ-OPP-2012-0641	Michael Glikes <i>glikes.michael@epa.gov</i> (703) 231-6499
Iodine and Iodophors Case 3080	EPA-HQ-OPP-2013-0767	Michael McCarroll <i>mccarroll.michael@epa.gov</i> (703) 347-0147
Metam Sodium and Metam Potassium Case 2390	EPA-HQ-OPP-2013-0140	Tiffany Green <i>green.tiffany@epa.gov</i> (703) 347-0314
Methyl isothiocyanate (MITC) Case 2405	EPA-HQ-OPP-2013-0242	Megan Snyderman <i>snyderman.megan@epa.gov</i> (703) 347-0671
Pelargonic Acid, Salts and Esters Case 6077	EPA-HQ-OPP-2010-0424	Michael McCarroll <i>mccarroll.michael@epa.gov</i> (703) 347-0147
Phenmedipham	EPA-HQ-OPP-2014-0546	Lauren Bailey

Case 0277		<i>bailey.lauren@epa.gov</i> (703) 347-0374
Pythium oligandrum DV 74 Case 6511	EPA-HQ-OPP-2017-0393	Cody Kendrick <i>kendrick.cody@epa.gov</i> (703) 347-0468
Sethoxydim Case 2600	EPA-HQ-OPP-2015-0088	Steven R. Peterson <i>peterson.stevenr@epa.gov</i> (703) 347-0755
Tetraacetythylenediamine (TAED) Case 5105	EPA-HQ-OPP-2013-0608	Kimberly Wilson <i>wilson.kimberly@epa.gov</i> (703) 347-0495
Thymol Case 3143	EPA-HQ-OPP-2010-0002	Kimberly Wilson <i>wilson.kimberly@epa.gov</i> (703) 347-0495
Tralopyril (Econea) Case 5114	EPA-HQ-OPP-2013-0217	Erin Dandridge <i>dandridge.erin@epa.gov</i> (703) 347-0185
Triclosan Case 2340	EPA-HQ-OPP-2012-0811	Megan Snyderman <i>snyderman.megan@epa.gov</i> (703) 347-0671

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: October 30, 2019.

**Mary Reaves,**

*Acting Director,*

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

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