TABLE 1—PROPOSED INTERIM DECISIONS—Continued

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Tetraconazole, Case Number 7043.	EPA–HQ–OPP–2015–0061	Veronica Dutch, *dutch.veronica@epa.gov,* (703) 308–8585.

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 18, 2021.

Mary Elissa Reaves,

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

[FR Doc. 2021–23043 Filed 10–21–21; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0751; FRL–9076–01–OCSPP]

Pesticide Registration Review; Interim Decisions and Case Closures for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's interim registration review decisions for the following chemicals: Amicarbazone; aminopyralid; azadirachtin, cold pressed neem oil and clarified hydrophobic neem oil; benzoic acid; endothall and salts; ethofumesate; fluoxastrobin; forchlorfenuron; gamma-cyhalothrin; inorganic halides; ipconazole; L-lactic acid; lambda-cyhalothrin; metam/MITC; metconazole; myclobutanil; novaluron; picloram; prometon; prothioconazole; and pyrasulfotole. In addition, it announces the closure of the registration review case for propazine.

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 pesticide specific contact person listed in the Table in Unit IV.

B. How can I get copies of this document and other related information?

The dockets these cases, identified by the docket identification (ID) number for the specific pesticide of interest provided in the Table in Unit IV., are available at *http://www.regulations.gov* or at the Office of Pesticide Programs Regulatory Public Docket (OPP Docket) in the Environmental Protection Agency Docket Center (EPA/DC), West William Jefferson Clinton Bldg., Rm. 3334, 1301 Constitution Ave. NW, Washington, DC 20460–0001. The Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566–1744, and the telephone number for the OPP Docket is (703) 305–5805.

Due to the public health concerns related to COVID–19, the EPA Docket Center (EPA/DC) and Reading Room is closed to visitors with limited exceptions. The staff continues to provide remote customer service via email, phone, and webform. For the latest status information on EPA/DC services and docket access, visit *https://www.epa.gov/dockets.*

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 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 interim registration review decisions for the pesticides shown in the following table. The interim registration review decisions are supported by rationales included in the docket established for each chemical.

TABLE—REGISTRATION REVIEW INTERIM DECISIONS BEING ISSUED

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Amicarbazone, Case Number 7262	EPA–HQ–OPP–2015–0400	Samantha Thomas, *thomas.samantha@epa.gov*, (703) 347–0514.
Aminopyralid, Case Number 7267	EPA–HQ–OPP–2013–0749	Rachel Stephenson, *stephenson.rachel@epa.gov*, (703) 347–8904.
Azadirachtin, Cold Pressed Neem Oil, and Clarified Hydrophobic Neem Oil, Case Number 6021.	EPA–HQ–OPP–2008–0632	Joseph Mabon, *mabon.joseph@epa.gov*, (703) 347–0177.
Benzoic Acid, Case Number 5107	EPA–HQ–OPP–2010–0692	Megan Snyderman, *snyderman.megan@epa.gov*, (703) 347–0671.
Endothall and Salts, Case Number 2245	EPA–HQ–OPP–2015–0591	Robert Little, *little.robert@epa.gov*, (703) 347–8156.
Ethofumesate, Case Number 2265	EPA–HQ–OPP–2015–0406	James Douglass, *douglass.james@epa.gov*, (703) 347–8630.
Fluoxastrobin, Case Number 7044	EPA–HQ–OPP–2015–0295	Rachel Fletcher, *fletcher.rachel@epa.gov*, (703) 347–0512.
Forchlorfenuron, Case Number 7057	EPA–HQ–OPP–2014–0641	Srijana Shrestha, *shrestha.srijana@epa.gov*, (703) 305–6471.
Gamma-cyhalothrin, Case Number 7437	EPA–HQ–OPP–2010–0479	Darius Stanton, *stanton.darius@epa.gov*, (703) 347–0433.
Inorganic halides, Case Number 4051	EPA–HQ–OPP–2009–0168	Erin Dandridge, *dandridge.erin@epa.gov*, (703) 347–0185.
Ipconazole, Case Number 7041	EPA–HQ–OPP–2015–0590	Alex Hazlehurst, *hazlehurst.alexander@epa.gov*, (703) 347–0221.
L-lactic Acid, Case Number 6062	EPA–HQ–OPP–2020–0552	SanYvette Williams, *williams.sanyvette@epa.gov*, (703) 305–7702.
Lambda-cyhalothrin, Case number 7408	EPA–HQ–OPP–2010–0480	Darius Stanton, *stanton.darius@epa.gov*, (703) 347–0433.
Metam/MITC, Case Number 2390 & 2405	EPA–HQ–OPP–2013–0140 & EPA–HQ–OPP–2013–0242	Tiffany Green, *green.tiffany@epa.gov*, (703) 347–0314.
Metconazole, Case Number 7049	EPA–HQ–OPP–2015–0013	Samantha Thomas, *thomas.samantha@epa.gov*, (703) 347–0514.
Myclobutanil, Case Number 7006	EPA–HQ–OPP–2015–0053	Anitha Kisanga, *kisanga.anitha@epa.gov*, (703) 347–0540.
Novaluron Case Number 7615	EPA–HQ–OPP–2015–0171	Robert Little, *little.robert@epa.gov*, (703) 347–8156.
Picloram, Case Number 0096	EPA–HQ–OPP–2013–0740	Andy Muench, *muench.andrew@epa.gov*, (703) 347–8263.
Prometon, Case Number 2545	EPA–HQ–OPP–2013–0068	Carolyn Smith, *smith.carolyn@epa.gov*, (703) 347–8325.
Prothioconazole, Case Number 7054	EPA–HQ–OPP–2015–0474	Rachel Eberius, *eberius.rachel@epa.gov*, (703) 347–0492.
Pyrasulfotole, Case Number 7272	EPA–HQ–OPP–2016–0391	James Douglass, *douglass.james@epa.gov*, (703) 347–8630.

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 propazine (Case Number 0230, Docket ID Number EPA–HQ–OPP–2013–0250) 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: October 18, 2021.

Mary Elissa Reaves,

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

[FR Doc. 2021–23041 Filed 10–21–21; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2021–0068; FRL–8732–04–OCSPP]

Certain New Chemicals; Receipt and Status Information for September 2021

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA is required under the Toxic Substances Control Act (TSCA) to make information publicly available and to publish information in the **Federal Register** pertaining to submissions under TSCA Section 5, including notice of receipt of a Premanufacture notice (PMN), Significant New Use Notice (SNUN) or Microbial Commercial Activity Notice (MCAN), including an amended notice or test information; an exemption application (Biotech exemption); an application for a test marketing exemption (TME), both pending and/or concluded; a notice of commencement (NOC) of manufacture (including import) for new chemical substances; and a periodic status report on new chemical substances that are currently under EPA review or have recently concluded review. This document covers the period from 09/01/2021 to 09/30/2021.

DATES: Comments identified by the specific case number provided in this document must be received on or before November 22, 2021.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPPT–2021–0068, and the specific case number for the chemical substance related to your comment, through the Federal eRulemaking Portal at *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. Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is available at *http://www.epa.gov/dockets.*

Due to the public health concerns related to COVID–19, the EPA Docket Center (EPA/DC) and Reading Room is closed to visitors with limited exceptions. The staff continues to provide remote customer service via email, phone, and webform. For the latest status information on EPA/DC services and docket access, visit *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: *For technical information contact:* Jim Rahai, Project Management and Operations Division (MC 7407M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 564–8593; email address: *rahai.jim@epa.gov*.

For general information contact: The TSCA-Hotline, ABVI-Goodwill, 422 South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554–1404; email address: *TSCA-Hotline@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. Executive Summary

A. What action is the Agency taking?

This document provides the receipt and status reports for the period from 09/01/2021 to 09/30/2021. The Agency is providing notice of receipt of PMNs, SNUNs and MCANs (including amended notices and test information); an exemption application under 40 CFR part 725 (Biotech exemption); TMEs, both pending and/or concluded; NOCs to manufacture a new chemical substance; and a periodic status report on new chemical substances that are currently under EPA review or have recently concluded review.

EPA is also providing information on its website about cases reviewed under the amended TSCA, including the section 5 PMN/SNUN/MCAN and exemption notices received, the date of receipt, the final EPA determination on the notice, and the effective date of EPA's determination for PMN/SNUN/MCAN notices on its website at: *https://www.epa.gov/reviewing-new-chemicals-under-toxic-substances-control-act-tsca/status-pre-manufacture-notices.* This information is updated on a weekly basis.

B. What is the Agency's authority for taking this action?

Under the Toxic Substances Control Act (TSCA), 15 U.S.C. 2601 *et seq.,* a chemical substance may be either an ''existing'' chemical substance or a ''new'' chemical substance. Any chemical substance that is not on EPA's TSCA Inventory of Chemical Substances (TSCA Inventory) is classified as a ''new chemical substance,'' while a chemical substance that is listed on the TSCA Inventory is classified as an ''existing chemical substance.'' (See TSCA section 3(11).) For more information about the TSCA Inventory please go to: *https://www.epa.gov/tsca-inventory.*

Any person who intends to manufacture (including import) a new chemical substance for a non-exempt commercial purpose, or to manufacture or process a chemical substance in a non-exempt manner for a use that EPA has determined is a significant new use, is required by TSCA section 5 to provide EPA with a PMN, MCAN or SNUN, as appropriate, before initiating the activity. EPA will review the notice, make a risk determination on the chemical substance or significant new use, and take appropriate action as described in TSCA section 5(a)(3).

TSCA section 5(h)(1) authorizes EPA to allow persons, upon application and under appropriate restrictions, to manufacture or process a new chemical substance, or a chemical substance subject to a significant new use rule (SNUR) issued under TSCA section 5(a)(2), for ''test marketing'' purposes, upon a showing that the manufacture, processing, distribution in commerce, use, and disposal of the chemical will not present an unreasonable risk of injury to health or the environment. This is referred to as a test marketing exemption, or TME. For more information about the requirements applicable to a new chemical go to: *http://www.epa.gov/oppt/newchems.*

Under TSCA sections 5 and 8 and EPA regulations, EPA is required to publish in the **Federal Register** certain information, including notice of receipt of a PMN/SNUN/MCAN (including amended notices and test information); an exemption application under 40 CFR part 725 (biotech exemption); an application for a TME, both pending and concluded; NOCs to manufacture a new chemical substance; and a periodic status report on the new chemical substances that are currently under EPA review or have recently concluded review.

C. Does this action apply to me?

This action provides information that is directed to the public in general.

D. Does this action have any incremental economic impacts or paperwork burdens?

No.

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

1. *Submitting confidential business information (CBI).* Do not submit this information to EPA through