

information about the docket available at <http://www.epa.gov/dockets>.

## II. Background

The Frank R. Lautenberg Chemical Safety for the 21st Century Act, amending the Toxic Substances Control Act of 1976, was signed into law on June 22, 2016. The amendments have enhanced EPA's authority to evaluate chemical substances.

Since the 2016 amendments to TSCA, EPA has been working to make the policy and process changes necessary to align the New Chemicals program with the requirements of the new law, as well as to streamline and improve the review process. In November 2017, EPA released the "*New Chemicals Decision-Making Framework: Working Approach to Making Determinations under section 5 of TSCA*" (the "Working Approach") for public comment, and subsequently held a public meeting on implementing the New Chemicals program under amended TSCA on December 14, 2017.

After consideration of comments received on the 2017 version and based on additional implementation experience, EPA is updating the Working Approach. Later in December 2019, EPA will announce the availability of the updated document after the public meeting and will accept comments on the updated document. EPA expects the updated document will provide further clarity and detail on EPA's approach and practices, including: (1) EPA's general guiding principles and concepts for making determinations on new chemical notices submitted to EPA under section 5 of TSCA; (2) the decision-making logic and the key questions that EPA must address; and (3) a discussion of how EPA might apply the working approach to reach one of the five new chemical determinations allowable under the statute.

Additional information on the TSCA amendments can be found at <https://www.epa.gov/assessing-and-managing-chemicals-under-tsca/frank-r-lautenberg-chemical-safety-21st-century-act>.

## III. Meeting

### A. Remote Access

The meeting will be accessible remotely for registered participants. Registered participants will receive information on how to connect to the meeting prior to its start.

### B. Public Participation at the Meeting

Members of the public may register to attend the meeting as observers and may also register to speak at the meeting,

using one of the registration methods described under **ADDRESSES**. A registered speaker is encouraged to focus on issues directly relevant to the meeting's subject matter. Each speaker will be allowed approximately three minutes to provide oral feedback, subject to the number of confirmed registered speakers. A speaker must be registered in order to speak during the meeting. To accommodate as many registered speakers as possible, speakers may not use visual aids or written material. Persons registered to speak (as well as others) may submit written materials to the dockets as described under **ADDRESSES**. The meeting agenda and supporting materials will be made available in the docket and on EPA's website in advance of the meeting.

## IV. How can I request to participate in these meeting?

### A. Registration

To attend the meeting in person or to receive remote access, you must register online no later than December 6, 2019, using one of the methods described under **ADDRESSES**. While on-site registration will be available, seating will be on a first-come, first-served basis, with priority given to early registrants, until room capacity is reached. For registrants not able to attend in person, the meeting will also provide remote access capabilities; registered participants will be provided information on how to connect to the meeting prior to its start.

### B. Required Registration Information

Members of the public may register to attend as observers or to speak during the scheduled public speaking period at the meeting. To register for the meeting online, you must provide your full name, organization or affiliation, and contact information.

**Authority:** 15 U.S.C. 2601 *et seq.*

Dated: November 15, 2019.

**Tala Henry,**

*Deputy Director, Office of Pollution Prevention and Toxics.*

[FR Doc. 2019-25171 Filed 11-19-19; 8:45 am]

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## ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2017-0720; FRL-10001-76]

### 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: Pyrimethanil and saflufenacil. With this document, the EPA is opening the public comment period for registration review for these chemicals. This notice also announces the availability of EPA's draft ecological risk assessment for the pesticide pyrimethanil and opens a 60-day public comment period on the draft risk assessment. EPA is also announcing that it will not be opening a docket, nor conducting registration review for meptyldinocap.

**DATES:** Comments must be received on or before January 21, 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:* ÖPP 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 the Table 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](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 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. This notice also announces the availability of EPA's draft ecological risk assessment for the pesticide pyrimethanil and opens a 60-day public comment period on the draft risk assessment.

Registration review case name and number	Docket ID No.	Chemical review manager and contact information
Pyrimethanil Case 7059 .....	EPA-HQ-OPP-2019-0380	Lauren Bailey, <a href="mailto:bailey.lauren@epa.gov">bailey.lauren@epa.gov</a> , (703) 347-0374.
Saflufenacil Case 7278 .....	EPA-HQ-OPP-2019-0524	Jonathan Williams, <a href="mailto:williams.jonathanr@epa.gov">williams.jonathanr@epa.gov</a> , (703) 347-0670.

EPA is also announcing that it will not be opening a docket, nor taking public comments for meptyldinocap. Meptyldinocap is a fungicide used on grapes abroad, and a tolerance was established in 2009 for residues of meptyldinocap on grapes imported into the United States. There are have been no changes since the initial tolerance decision and no changes to the tolerance or tolerance definition are being proposed. Meptyldinocap does not have any products registered in the United States under FIFRA section 3, therefore meptyldinocap is not scheduled for review under the registration review program.

*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: October 29, 2019.

**Mary Reaves,**

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

[FR Doc. 2019-25164 Filed 11-19-19; 8:45 am]

**BILLING CODE 6560-50-P**

## FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060-0768]

### Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice and request for comments.

**SUMMARY:** As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act of 1995 (PRA), the Federal Communications Commission (FCC or Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid Office of Management and Budget (OMB) control

number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

**DATES:** Written PRA comments should be submitted on or before January 21, 2020. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

**ADDRESSES:** Direct all PRA comments to Cathy Williams, FCC, via email [PRA@fcc.gov](mailto:PRA@fcc.gov) and to [Cathy.Williams@fcc.gov](mailto:Cathy.Williams@fcc.gov).

**FOR FURTHER INFORMATION CONTACT:** For additional information about the information collection, contact Cathy Williams, (202) 418-2918.

**SUPPLEMENTARY INFORMATION:**

*OMB Control No.:* 3060-0768.

*Title:* 28 GHz Band Segmentation Plan Amending the Commission's Rules to Redesignate the 27.5-29.5 GHz Frequency Band, to Reallocate the 29.5 to 30.0 GHz Frequency Band and to Establish Rules and Policies.

*Form No.:* None.

*Type of Review:* Extension of a currently approved collection.

*Respondents:* Business or other for-profit entities.

*Number of Respondents/Responses:* 17 respondents; 17 responses.

*Estimated Time per Response:* 2 hours.

*Frequency of Response:* On occasion reporting requirement; third-party disclosure requirement.

*Obligation to Respond:* Required to obtain or retain benefits. The statutory authority for this information collection is contained in 47 U.S.C. 154 and 303.

*Total Annual Burden:* 34 hours.

*Annual Cost Burden:* \$4,950.

*Privacy Act Impact Assessment:* No impact(s).

*Nature and Extent of Confidentiality:* In general, there is no need for confidentiality with this collection of information.

*Needs and Uses:* The Federal Communications Commission ("Commission") is requesting an extension of the information collection titled, "28 GHz Band" under OMB Control No. 3060-0768 from the Office of Management and Budget (OMB).

The information collection requirements contained in this collection require are as follows: (1) Local Multipoint Distribution Systems (LMDS) licensees to serve copies of their applications on all Non-Geostationary Mobile Satellite Service (NGSO/MSS) applicants (Section 101.147) and (2) NGSO/MSS feeder link earth stations

must specify a set of geographic coordinates for location of these earth stations, 15 days after the release of a public notice announcing commencement of LMDS auctions (Section 101.147).

The information is used by the Commission and other applicants and/or licensees in the 28 GHz band to facilitate technical coordination of systems among applicants and/or licensees in the 28 GHz band. Without such information, the Commission could not implement the Commission's band plan. Affected applicants and licensees are required to provide the requested information to the Commission and other third parties whenever they seek authority to provide service in the 28 GHz band. The frequency of filing is, in general, determined by the applicant or licensees. If this information is compiled less frequently or not filed in conjunction with our rules, applicants and licensees will not obtain the authorization necessary to provide telecommunications services. Furthermore, the Commission would not be able to carry out its mandate as required by statute and applicants and licensees would not be able to provide service effectively.

Federal Communications Commission.

**Marlene Dortch,**

*Secretary, Office of the Secretary.*

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## FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060-0398]

### Information Collection Being Submitted for Review and Approval to the Office of Management and Budget

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice and request for comments.

**SUMMARY:** As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (PRA) of 1995, the Federal Communications Commission (FCC or the Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collection. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility;