contact OPP at (202) 502–6595 or *OPP@ferc.gov*.

Dated: April 24, 2024.

Debbie-Anne A. Reese,

Acting Secretary.

[FR Doc. 2024–09290 Filed 4–29–24; 8:45 am]

BILLING CODE 6717–01–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0750; FRL–11907–01–OCSPP]

Pesticide Registration Review; Proposed 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 decisions (PIDs) and amended PIDs for the following pesticides: Acephate, Captan, Ferbam, Thiram, and Ziram. EPA is opening a 60-day public comment period for these proposed interim registration review decisions.

DATES: Comments must be received on or before July 1, 2024.

ADDRESSES: Submit your comments through *https://www.regulations.gov* using the docket identification (ID) number for the pesticide of interest as identified in Table 1 of Unit I. 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 and visiting the docket, along with more information about dockets generally, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT:

For pesticide specific information: The Chemical Review Manager for the pesticide of interest is identified in Table 1 of Unit I.

For general information: Melanie Biscoe, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–0701; email address: *biscoe.melanie@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. What action is the Agency taking?

Pursuant to 40 CFR 155.58(a), this notice announces the availability of EPA's proposed interim and proposed 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 REGISTRATION REVIEW DECISIONS

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Acephate Case Number 0042	EPA–HQ–OPP–2008–0915.	Kent Fothergill, *fothergill.kent@epa.gov,* (202) 566–1943.
Captan (Amended) Case Number 0120	EPA–HQ–OPP–2013–0296.	Christina Scheltema, *scheltema.christina@epa.gov,* (202) 566–2272.
Ferbam (Amended) Case Number 8000	EPA–HQ–OPP–2015–0567.	DeMariah Koger, *koger.demariah@epa.gov,* (202) 566–2288.
Thiram (Amended) Case Number 0122	EPA–HQ–OPP–2015–0433.	DeMariah Koger, *koger.demariah@epa.gov,* (202) 566–2288.
Ziram (Amended) Case Number 8001	EPA–HQ–OPP–2015–0568.	DeMariah Koger, *koger.demariah@epa.gov,* (202) 566–2288.

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

EPA is conducting its registration review of the chemicals listed in the table 1 of unit I pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) section 3(g) (7 U.S.C. 136a(g)) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. FIFRA section 3(g) provides, among other things, that pesticide registrations are to be reviewed every 15 years. Consistent with 40 CFR 155.57, in its final registration review decision, EPA will ultimately determine whether a pesticide continues to meet the registration standard in FIFRA section 3(c)(5) (7 U.S.C. 136a(c)(5)). As part of the registration review process, the Agency has completed a proposed interim or proposed decision for each of the pesticides listed in Table 1 of Unit I.

The registration review docket for a pesticide includes documents related to the registration review case. Among other things, these documents describe EPA's rationales for conducting additional risk assessments for the registration review of the pesticides included in Table 1 of Unit I, as well as the Agency's subsequent risk findings and consideration of possible risk mitigation measures. The proposed interim and proposed registration review decisions are supported by the rationales included in those documents.

Consistent with 40 CFR 155.58(a), EPA provides for at least a 60-day public comment period on proposed interim and proposed 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 decision.

For additional background on the registration review program, see: *https://www.epa.gov/pesticide-reevaluation.*

III. Does this action apply to me?

This notice 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 table 1 of unit I.

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

In submitting a comment to EPA, please consider the following:

1. *Submitting CBI.* Do not submit this information to EPA through *regulations.gov* or email. Clearly mark the part or all 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: *https://www.epa.gov/dockets/commenting-epa-dockets.*

3. *Environmental justice.* EPA seeks to achieve environmental justice, the fair treatment and meaningful involvement of any group, including minority and/or low-income populations, in the development, implementation, and enforcement of environmental laws, regulations, and policies. To help address potential environmental justice issues, the Agency seeks information on any groups or segments of the population who, as a result of their location, cultural practices, or other factors, may have atypical or disproportionately high and adverse human health impacts or environmental effects from exposure to the pesticides discussed in this document, compared to the general population.

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 Table 1 in Unit I. The Agency will consider all comments received by the closing date and may respond to comments in a ''Response to Comments Memorandum'' in the docket and/or in any subsequent interim or final registration review decision, as appropriate.

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

Dated: April 23, 2024.

Timothy Kiely,

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

[FR Doc. 2024–09181 Filed 4–29–24; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0636; FR ID 216233]

Information Collection Being Reviewed by the Federal Communications Commission

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; 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.

DATES: Written PRA comments should be submitted on or before July 1, 2024. 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 Nicole Ongele, FCC, via email *PRA@fcc.gov* and to *nicole.ongele@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Nicole Ongele, (202) 418–2991.

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

OMB Control Number: 3060–0636.

Title: Sections 2.906, 2.909, 2.1071, 2.1074, 2.1077 and 15.37, Equipment Authorizations—Supplier's Declaration of Conformity (SDoC).

Form No.: N/A.

Type of Review: Revision of a currently approved collection.

Respondents: Business or other for-profit entities.

Number of Respondents and Responses: 8,500 respondents; 17,000 responses.

Estimated Time per Response: 1–18 hours (average).

Frequency of Response: One-time reporting requirement, recordkeeping requirement and third party disclosure requirements.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. 154(i), 301, 302a, 303, 309(j), 312, 403, 503, and the Secure Equipment Act of 2021, Public Law 117–55, 135 Stat. 423.

Total Annual Burden: 161,500 hours.

Total Annual Cost: $17,000,000.

Needs and Uses: The Commission will submit this revised information collection to the Office of Management and Budget (OMB) after this 60 day comment period in order to obtain the full three year clearance from them.

In 2022, the Supplier's Declaration of Conformity (SDOC) procedure were revised in a Report and Order, FCC 22–84 (88 FR 7592, February 6, 2023). Revisions to the information collection included amendments to rule sections 2.906 and 2.909 as reported herein, therefore, the eligibility restrictions resulted in fewer applicants but the continued growth in participation in the program resulted in a re-adjustment of applicants which supports program changes and adjustments.

§ 2.906 Supplier's Declaration of Conformity

(a) Supplier's Declaration of Conformity (SDoC) is a procedure where the responsible party, as defined in § 2.909, makes measurements or completes other procedures found acceptable to the Commission to ensure that the equipment complies with the appropriate technical standards and other applicable requirements. Submittal to the Commission of a sample unit or representative data demonstrating compliance is not required unless specifically requested pursuant to § 2.945.

(b) Supplier's Declaration of Conformity is applicable to all items subsequently marketed by the manufacturer, importer, or the responsible party that are identical, as defined in § 2.908, to the sample tested and found acceptable by the manufacturer.

(c) The responsible party may, if it desires, apply for Certification of a device subject to the Supplier's Declaration of Conformity. In such cases, all rules governing certification will apply to that device.

(d) Notwithstanding other parts of this section, equipment otherwise subject to the Supplier's Declaration of Conformity process that is produced by any entity