upfront substantiation requirement. Lastly, EPA agrees that under 40 CFR 158 section 158.33(d)(1), information properly designated by the submitter as FIFRA section 10(d)(1)(A), (B), or (C) may be considered for confidential protection; however, as discussed in the draft PR notice, none of the information provided on EPA Form No. 8570–35 is within the scope of FIFRA section 10(d)(1) (A), (B), or (C).

III. Do PR Notices contain binding requirements?

The PR Notice discussed in this document is intended to provide guidance to EPA personnel and decision makers and to pesticide registrants. While the requirements in the statutes and Agency regulations are binding on EPA and the applicants, this PR Notice is not binding on either EPA or pesticide registrants, and EPA may depart from the guidance where circumstances warrant and without prior notice. Likewise, pesticide registrants may assert that the guidance is not appropriate generally or not applicable to a specific pesticide or situation.

IV. Are these forms approved under the Paperwork Reduction Act (PRA)?

According to the PRA, 44 U.S.C. 3501 *et seq.,* an agency may not conduct or sponsor, and a person is not required to respond to a collection of information that requires approval under the PRA, unless it has been approved by the Office of Management and Budget (OMB) and displays a currently valid OMB control number. The information collection activities associated with EPA Forms 8570–34 and 8570–35 and the activities described in this PR Notice are already approved by OMB under the PRA and are contained in the Information Collection Requests (ICRs) entitled ''Consolidated Pesticide Registration Submission Portal'' identified as EPA ICR No. 2624.01 and approved under OMB Control No. 2070–0226; and the ''Pesticide Data Call-in Program'' identified as EPA ICR No. 2288 and approved under OMB Control No. 2070–0174. For additional information about these ICRs, go to *https://www.reginfo.gov/public/* and search on the applicable OMB control number.

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

Dated: March 28, 2025.

Edward Messina,

Director, Office of Pesticide Programs.

[FR Doc. 2025–05866 Filed 4–3–25; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2025–0077; FRL–12476–01–OCSPP]

Certain New Chemicals or Significant New Uses; Statements of Findings for January 2025

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Toxic Substances Control Act (TSCA) requires EPA to publish in the **Federal Register** a statement of its findings after its review of certain TSCA submissions when EPA makes a finding that a new chemical substance or significant new use is not likely to present an unreasonable risk of injury to health or the environment. Such statements apply to premanufacture notices (PMNs), microbial commercial activity notices (MCANs), and significant new use notices (SNUNs) submitted to EPA under TSCA. This document presents statements of findings made by EPA on such submissions during the period from January 1, 2025 to January 31, 2025. This document also presents statements of findings on submissions made by EPA during earlier time periods that were inadvertently omitted from notices for those time periods that were identified during a quality control review.

ADDRESSES: The docket for this action, identified by docket identification (ID) number EPA–HQ–OPPT–2025–0077, is available online at *https://www.regulations.gov.* Additional information about dockets generally, along with instructions for visiting the docket in-person, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT:

For technical information contact: Rebecca Edelstein, New Chemical Division (7405M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 564–1667; email address: *edelstein.rebecca@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. Does this action apply to me?

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

B. What action is the Agency taking?

This document lists the statements of findings made by EPA after review of submissions under TSCA section 5(a) that certain new chemical substances or significant new uses are not likely to present an unreasonable risk of injury to health or the environment. This document presents statements of findings made by EPA during the reporting period, and several made by EPA during earlier time periods that were inadvertently omitted from notices for those time periods that were identified during a quality control review.

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

TSCA section 5(a)(3) requires EPA to review a submission under TSCA section 5(a) and make one of several specific findings pertaining to whether the substance may present unreasonable risk of injury to health or the environment. Among those potential findings is that the chemical substance or significant new use is not likely to present an unreasonable risk of injury to health or the environment per TSCA Section 5(a)(3)(C).

TSCA section 5(g) requires EPA to publish in the **Federal Register** a statement of its findings after its review of a submission under TSCA section 5(a) when EPA makes a finding that a new chemical substance or significant new use is not likely to present an unreasonable risk of injury to health or the environment. Such statements apply to PMNs, MCANs, and SNUNs submitted to EPA under TSCA section 5.

Anyone who plans to manufacture (which includes import) a new chemical substance for a non-exempt commercial purpose and any manufacturer or processor wishing to engage in a use of a chemical substance designated by EPA as a significant new use must submit a notice to EPA at least 90 days before commencing manufacture of the new chemical substance or before engaging in the significant new use.

The submitter of a notice to EPA for which EPA has made a finding of ''not likely to present an unreasonable risk of injury to health or the environment'' may commence manufacture of the chemical substance or manufacture or processing for the significant new use notwithstanding any remaining portion of the applicable review period.

II. Statements of Findings Under TSCA Section 5(a)(3)(C)

In this unit, EPA identifies the PMNs, MCANs and SNUNs for which EPA has made findings under TSCA section 5(a)(3)(C) that the new chemical substances or significant new uses are not likely to present an unreasonable risk of injury to health or the environment.

For the findings made during this period, the following list provides the EPA case number assigned to the TSCA section 5(a) submission and the chemical identity (generic name if the specific name is claimed as Confidential Business Information (CBI)), and the date the findings were made:

• P–24–0124, Alkyl alkenoic acid, alkyl ester, polymer with substituted carbomonocycle, substituted alkyl alkyl alkenoate phosphate, substituted alkyl alkyl alkenoate, alkyl alkenoate, heteromonocycle polymer, substitutes alkenyl substituted alkyl ester, alkyl substituted alkyl alkanoate initiated (Generic Name); Decision date: 01/29/2025.

In addition, the following list provides the same information for those cases where EPA made the findings during earlier time periods that were inadvertently omitted from notices for those time periods as identified during a quality control review:

• J–19–0026–0027, Biofuel-producing modified microorganism(s), with chromosomally-borne modifications (Generic Name). Decision date: 01/23/2020.

• J–23–0005–0006, Microorganisms genetically transformed to express an enzyme (Generic Name). Decision date: 11/27/2023.

• P–23–0028, Gelatin and maltodextrin crosslinked with linear and cyclic aliphatic polyisocyanates (Generic Name). Decision date: 12/14/2023.

• P–23–0101, Glycerides from fermentation of genetically modified microorganism, epoxidized (Generic Name). Decision date: 11/13/2024.

• P–23–0102, Glycerides from fermentation of genetically modified microorganism (Generic Name). Decision date: 06/07/2024.

• P–23–0103, Glycerides from fermentation of genetically modified microorganism, epoxidized, reaction products with ethanol (Generic Name). Decision date: 11/13/2024.

• P–23–0143, L-Lysine, N-(3-carboxy-1-oxopropyl) derivs., sodium salts; CASRN 1917323–94–4. Decision date: 08/22/2024.

• P–24–0005, [1,1′-Biphenyl]-4,4′-diol, reaction products with 4-cyclohexylphenol and 2,4,6-trichloro-1,3,5-triazine; CASRN 2561414–35–3. Decision date: 07/16/2024.

• P–24–0106–0107, Glycerides, alkyl, epoxidized, alkyl esters (Generic Name). Decision date: 12/19/2024.

To access EPA's decision document describing the basis of the ''not likely to present an unreasonable risk'' finding made by EPA under TSCA section 5(a)(3)(C), look up the specific case number at *https://www.epa.gov/reviewing-new-chemicals-under-toxic-substances-control-act-tsca/chemicals-determined-not-likely.*

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

Dated: March 25, 2025.

Shari Z. Barash,

Director, New Chemicals Division, Office of Pollution Prevention and Toxics.

[FR Doc. 2025–05821 Filed 4–3–25; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0010; FR ID 287162]

Information Collection Being Submitted for Review and Approval to 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, 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. Pursuant to the Small Business Paperwork Relief Act of 2002, the FCC seeks specific comment on how it might ''further reduce the information collection burden for small business concerns with fewer than 25 employees.''

DATES: Written comments and recommendations for the proposed information collection should be submitted on or before May 5, 2025.

ADDRESSES: Comments should be sent to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. Your comment must be submitted into *www.reginfo.gov* per the above instructions for it to be considered. In addition to submitting in *www.reginfo.gov* also send a copy of your comment on the proposed information collection to Cathy Williams, FCC, via email to *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.* Include in the comments the OMB control number as shown in the **SUPPLEMENTARY INFORMATION** below.

FOR FURTHER INFORMATION CONTACT: For additional information or copies of the information collection, contact Cathy Williams at (202) 418–2918. To view a copy of this information collection request (ICR) submitted to OMB: (1) go to the web page *http://www.reginfo.gov/public/do/PRAMain,* (2) look for the section of the web page called ''Currently Under Review,'' (3) click on the downward-pointing arrow in the ''Select Agency'' box below the ''Currently Under Review'' heading, (4) select ''Federal Communications Commission'' from the list of agencies presented in the ''Select Agency'' box, (5) click the ''Submit'' button to the right of the ''Select Agency'' box, (6) when the list of FCC ICRs currently under review appears, look for the Title of this ICR and then click on the ICR Reference Number. A copy of the FCC submission to OMB will be displayed.

SUPPLEMENTARY INFORMATION: The Commission 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.

As part of its continuing effort to reduce paperwork burdens, as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501–3520), the FCC invited the general public and other Federal Agencies to take this opportunity to comment on the following information collection. Comments are requested concerning: (a) 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; (b) the accuracy of the Commission's burden estimates; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) 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. Pursuant to the Small Business Paperwork Relief Act of 2002, Public Law 107–198, see 44 U.S.C. 3506(c)(4), the FCC seeks specific comment on how it might ''further