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 applicable period.

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 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).

- P–25–0085–0088, Arylcarboxylic acid, polymer with alkanol, ester with substituted polyalkylene glycol (P–25–0085) (Generic Name); polymer of arylcarboxylic acid, glycol, polyalkylene glycol ether (P–25–0086) (Generic Name); polymer of arylcarboxylic acid, alkanediol and alcohols, alkoxylated (P–25–0087) (Generic Name); and polymer of arylcarboxylic acid, glycols and polyalkylene glycol ether (P–25–0088) (Generic Name).
- P–25–0089, Polymer of arylcarboxylic acid, sulfoarylcarboxylic acid, alkyl ester salt, glycols and oligoethylene glycol ether (Generic Name).
- P–25–0104, Alkyl alkenoic acid, substituted heteropolycyclic substituted alkyl ester, polymer with substituted carbopolycyclic, alkyl alkenoate, alkyl carbopolycyclic alkyl alkenoate and heteromonocyclic alkyl alkenoate (Generic Name).
- P–25–0142, Propenoic acid, methyl-, [bis[dimethyl[-trimethyl-bis[trimethylsilyl)oxy]- disiloxanyl] ethyl] silyl] oxy]—dimethyl-[[trimethyl-bis[trimethylsilyl)oxy]-disiloxanyl] ethyl]-disiloxanyl] propyl ester (Generic Name).

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), lookup the specific case number at *https://www.epa.gov/reviewing-new-chemicals-under-toxic-substances-control-act-tsca/determined-not-likely.*

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

Dated: May 21, 2026.

Shari Z. Barash,

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

[FR Doc. 2026–10708 Filed 5–28–26; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2026–3532; FRL–13370–01–OCSPP]

Pesticide Experimental Use Permit; Receipt of Application; Comment Request (March 2026)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces EPA's receipt of an application 100506–EUP–R from the State University of New York College of Environmental Science and Forestry (SUNY ESF) requesting an experimental use permit (EUP) for OxO American Chestnut (oxalate oxidase OxO) and the genetic material necessary for its expression in American chestnut (*Castanea dentata*). The Agency has determined that the permit may be of regional and national significance. Therefore, because of the potential significance, EPA is seeking comments on this application.

DATES: Comments must be received on or before June 29, 2026.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2026–3532, through the *Federal eRulemaking Portal* at *https://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 on and visiting the docket, along with more information about dockets generally, are available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Each application summary in Unit II. specifies a contact division. The appropriate division contacts are identified as follows:

- BPPD (Biopesticides and Pollution Prevention Division) (Mail Code 7511M); Shannon Borges; main telephone number: (202) 566–1400; email address: *BPPDFRNotices@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

This action is directed to the public in general. Although this action may be of particular interest to those people who conduct or sponsor research on pesticides, the Agency has not attempted to describe all the specific entities that may be affected by this action.

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 in 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 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.*

II. What Action is the Agency Taking?

Under section 5 of the Federal Insecticide, Fungicide and Rodenticide Act (FIFRA), 7 U.S.C. 136c, EPA can allow manufacturers to field test pesticides under development. Manufacturers are required to obtain an EUP before testing new pesticides or new uses of pesticides if they conduct experimental field tests on 10 acres or more of land or one acre or more of water.

Pursuant to 40 CFR 172.11(a), the Agency has determined that the following EUP application may be of regional and national significance, and therefore is seeking public comment on the EUP application:

Experimental Use Permit Number: 100506–EUP–R. *Docket ID Number:* EPA–HQ–OPP–2026–3532. *Submitter:* State University of New York College of Environmental Science and Forestry (SUNY ESF), 1 Forestry Drive, Syracuse, NY 13210. *Pesticide Chemical:* OxO American Chestnut (oxalate oxidase OxO) and the genetic material necessary for its expression in American chestnut (*Castanea dentata*). *Summary of Request:* The State University of New York College of Environmental Science and Forestry is requesting an experimental use permit for the active ingredient OxO American Chestnut (oxalate oxidase OxO) and the genetic material necessary for its expression in American chestnut (*Castanea dentata*) for three years in Alabama, Arkansas, Connecticut, Delaware, District of Columbia (Washington, DC), Florida, Georgia, Illinois, Indiana, Kentucky, Maine, Maryland, Massachusetts, Michigan, Mississippi, Missouri, New Hampshire, New Jersey, New York, North Carolina, Ohio, Pennsylvania, Rhode Island, South Carolina, Tennessee, Vermont, Virginia, West Virginia, and Wisconsin, with a total area of up to 280 acres across all states. The OxO gene enables the American chestnut trees to detoxify oxalic acid produced by the fungal pathogen *Cryphonectria parasitica,* thereby allowing coexistence with the pathogen. The purpose of this EUP is to evaluate survival, early growth, and establishment success of transgenic OxO American chestnuts, compared with wild-type American chestnuts and/or other control trees, across a wide range of environmental conditions and planting scenarios. This testing will inform restoration management strategies across the native American chestnut range. *Date of Receipt:* February 27, 2026. *Contact:* BPPD.

Following the review of the application and any comments and data received in response to this solicitation, EPA will decide whether to issue or deny the EUP request, and if issued, the conditions under which it is to be conducted. Any issuance of an EUP will be announced in the **Federal Register**.

Authority: 21 U.S.C. 346a.

Dated: May 26, 2026.

Edward Messina,

Director, Office of Pesticide Programs.

[FR Doc. 2026–10714 Filed 5–28–26; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0249; FR ID 348748]

Information Collection Being Reviewed by the Federal Communications Commission

AGENCY: Federal Communications Commission.

ACTION: Notice; request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (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.

DATES: Written comments should be submitted on or before July 28, 2026. 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 contacts below as soon as possible.

ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.*

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

SUPPLEMENTARY INFORMATION: 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.

As part of its continuing effort to reduce paperwork burdens, and as required by the PRA of 1995 (44 U.S.C. 3501–3520), the FCC 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.

OMB Control Number: 3060–0249.

Title: Sections 74.781, 74.1281 and 78.69, Station Records.

Form Number: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business and other for-profit entities; not-for-profit institutions; State, Federal or Tribal governments.

Number of Respondents and Responses: 14,052 respondents; 19,077 responses.

Estimated Time per Response: 0.375 hour-1 hour.

Frequency of Response: Recordkeeping requirement.