copy of the proposed settlement agreement. The official public docket is available for public viewing at the EPA Docket Center, EPA West, Room 3334, 1301 Constitution Ave. NW, Washington, DC 20460. The EPA Docket Center 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.

The electronic version of the public docket for this action contains a copy of the proposed settlement agreement and is available through *https://regulations.gov.* You may use *https://www.regulations.gov* to submit or view public comments, access the index listing of the contents to the official public docket, and access those documents in the public docket that are available electronically. Once in the system, key in the appropriate docket identification number then select ''search''.

II. Additional Information About the Proposed Settlement Agreement

On May 4, 2020, Natural Resources Defense Council (NRDC) petitioned EPA, under the Federal Food, Drug, and Cosmetic Act (FFDCA), to revoke all tolerances for residues of five neonics on or in food—acetamiprid, clothianidin, dinotefuran, imidacloprid, and thiamethoxam. Plaintiffs subsequently filed a mandamus petition on October 29, 2025, alleging that EPA's failure to issue a final decision on the FFDCA petition constitutes an unreasonable delay under the All Writs Act, 28 U.S.C. 1651(a), and the Administrative Procedure Act (''APA''), 5 U.S.C. 555(b).

The proposed settlement agreement states that no later than April 30, 2027, EPA must issue a final decision on Petitioner's administrative petition under 21 U.S.C. 346a(d)(4)(A). Further, the proposed settlement agreement states that within 15 days of the execution of the proposed settlement agreement, the parties will file a joint motion to continue holding the litigation in abeyance pending the April 30, 2027 deadline for EPA's issuance of a final decision on the administrative petition.

For a period of thirty (30) days following the date of publication of this publication, the Agency will accept written comments relating to the proposed settlement agreement from persons who are not named as parties to the litigation in question. EPA or the Department of Justice may withdraw or withhold consent to the proposed settlement agreement if the comments disclose facts or considerations that indicate that such consent is inappropriate, improper, inadequate, or inconsistent with the requirements of the APA or FFDCA. Unless EPA or the Department of Justice determines that consent should be withdrawn, the terms of the proposed settlement agreement will be affirmed.

III. Additional Information About Commenting on the Proposed Settlement Agreement

Submit your comments, identified by Docket ID No. EPA–HQ–OGC–2026–6172 via *https://www.regulations.gov.* Once submitted, comments cannot be edited or removed from this docket. EPA may publish any comment received to its docket. Do not submit to EPA's docket at *https://www.regulations.gov* any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.,* on the web, cloud, or other file sharing system). For additional submission methods, the full public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit *https://www.epa.gov/dockets/commenting-epa-dockets.* For additional information about submitting information identified as CBI, please contact the person listed in the **FOR FURTHER INFORMATION CONTACT** section of this document. Note that written comments containing CBI and submitted by mail may be delayed and deliveries or couriers will be received by scheduled appointment only.

If you submit an electronic comment, EPA recommends that you include your name, mailing address, and an email or other contact information in the body of your comment. This ensures that you can be identified as the submitter of the comment and allows EPA to contact you if EPA cannot read your comment due to technical difficulties or needs further information on the substance of your comment. Any identifying or contact information provided in the body of a comment will be included as part of the comment that is placed in the official public docket and made available in EPA's electronic public docket. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment.

Use of the *http://www.regulations.gov* website to submit comments to EPA electronically is EPA's preferred method for receiving comments. The electronic public docket system is an ''anonymous access'' system, which means EPA will not know your identity, email address, or other contact information unless you provide it in the body of your comment.

Please ensure that your comments are submitted within the specified comment period. Comments received after the close of the comment period will be marked ''late.'' EPA is not required to consider these late comments.

Dated: August 13, 2026.

Randolph L. Hill,

Associate General Counsel.

[FR Doc. 2026–16938 Filed 8–19–26; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2026–0333; FRL–13200–06–OCSPP]

Pesticide Product Registration; Receipt of Applications for New Active Ingredients (June 2026)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of receipt and request for comment.

SUMMARY: This document announces the Agency's receipt of and solicits comments on applications to register pesticide products containing active ingredients not included in any currently registered pesticide products. The Agency is providing this notice in accordance with the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). EPA uses the month and year in the title to identify when the Agency compiled the applications identified in this notice of receipt Unit II. of this document identifies certain applications received in 2025 that are currently being evaluated by EPA, along with information about each application, including when it was received, who submitted the application, and the purpose of the application.

DATES: Comments must be received on or before September 21, 2026.

ADDRESSES: Submit your comments, identified by the docket identification (ID) number and the *EPA File Symbol* or the *EPA Registration Number* of interest as shown in Unit II. of this document, online 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:

- AD (Antimicrobials Division) (Mail Code 7510M); Kristin Willis, main telephone number: (202) 566–0793; email address: *ADFRNotices@epa.gov.*
- BPPD (Biopesticides and Pollution Prevention Division) (Mail Code 7511M); Shannon Borges; main telephone number: (202) 566–1400; email address: *BPPDFRNotices@epa.gov.*
- RD (Registration Division) (Mail Code 7505T); Charles Smith; main telephone number: (202) 566–1030; email address: *RDFRNotices@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 is the Agency's authority for taking this action?

EPA is taking this action pursuant to section 3(c)(4) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), 7 U.S.C. 136a(c)(4), and 40 CFR 152.102.

C. What action is the Agency taking?

EPA is hereby providing notice of receipt and opportunity to comment on applications to register pesticide products containing active ingredients not included in any currently registered pesticide products. Notice of receipt of these applications does not imply a decision by the Agency on these applications. The applications identified in this document were received since the last notice that was issued and are currently being evaluated by EPA in accordance with the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). For actions being evaluated under EPA's public participation process for registration actions, there will be an additional opportunity for public comment on the proposed decisions. Please see EPA's public participation website for additional information on this process (*https://www.epa.gov/pesticide-registration/public-participation-process-registration-actions*).

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

1. *Submitting CBI.* Do not submit CBI to EPA through *https://www.regulations.gov* or email. If you wish to include CBI in your comment, please follow the applicable instructions at *https://www.epa.gov/dockets/commenting-epa-dockets#rules* and clearly mark the information that you claim to be CBI. In addition to one complete version of the comment that includes CBI, a copy of the comment without CBI must be submitted for inclusion in the public docket. Information marked as CBI 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. Registration Applications Received

This unit provides the following information about the applications received: The EPA File Symbol or Registration number(s); EPA docket ID number for the application; Name and address of the applicant; Name of the active ingredient, product type and proposed uses; and the division to contact for that application. Additional information about the application may also be available in the docket for the application as identified in this unit.

- *EPA File Symbols:* 33906–GR; 33906–GN. *Docket ID number:* EPA–HQ–OPP–2026–1784. *Applicant:* Nissan Chemical Corporation, 5–1, Nihonbashi 2-Chome, Chuo-ku, Tokyo 103–6119, Japan. *Product names:* Iptriazopyrid Technical; Iptriazopyrid 100 g/L SC. *Active ingredient:* Herbicide—Iptriazopyrid at 98.5%; 9.1%. *Proposed use:* Rice. *Date of receipt:* November 26, 2025. *Contact:* RD.
- *EPA File Symbol:* 52991–LR. *Docket ID number:* EPA–HQ–OPP–2026–5413. *Applicant:* Bedoukian Research, Inc., 6 Commerce Drive, Danbury, CT 06810. *Product name:* Bedoukiab z-9-Hexadecenal Technical Pheromone. *Active ingredient:* SCLP pheromone—z-9-Hexadecenal at 90%. *Proposed use:* Mating Disruptor (used as an attractant for incorporation into end-use products, intended to control lepidopteran species). *Date of receipt:* October 21, 2025. *Contact:* BPPD.
- *EPA File Symbol:* 97144–U. *Docket ID number:* EPA–HQ–OPP–2026–4786. *Applicant:* ReliOx Corporation, 8475 Western Way, Suite 155, Jacksonville, FL 32256. *Product name:* Whiff!. *Active ingredient:* Styrene, divinylbenzene and ethyl styrene copolymer, chloromethyl trimethylamine (26.148%) as an activator for sodium chlorite to generate chlorine dioxide. *Product type:* Antimicrobial Pesticide. *Proposed uses:* Non-food use, indoor disinfectant deodorizer, bactericide and virucide for residential and commercial use. *Date of receipt:* May 20, 2025. *Contact:* AD.

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

Dated: August 13, 2026.

Elizabeth Vizard,

Acting Director, Office of Pesticide Programs.

[FR Doc. 2026–16974 Filed 8–19–26; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–1272; FR ID 362738]

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

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) of 1995, the Federal Communications Commission (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 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 Commission 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.