IV. Procedures for Withdrawal of Request

Registrants who choose to withdraw a request for product cancellation should submit the withdrawal in writing to the person listed under **FOR FURTHER INFORMATION CONTACT**. If the products have been subject to a previous cancellation action, the effective date of cancellation and all other provisions of any earlier cancellation action are controlling.

V. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products that are currently in the United States and that were packaged, labeled, and released for shipment prior to the effective date of the cancellation action.

In any order issued in response to these requests for cancellation of product registrations EPA proposes to include the following provisions for the treatment of any existing stocks of the products listed in Table 1 of Unit II.

For voluntary product cancellations, registrants will be permitted to sell and distribute existing stocks of voluntarily canceled products for 1 year after the effective date of the cancellation, which will be the date of publication of the cancellation order in the **Federal Register**. Thereafter, registrants will be prohibited from selling or distributing the products identified in Table 1 of Unit II., except for export consistent with FIFRA section 17 (7 U.S.C. 136o) or for proper disposal.

Persons other than the registrant may sell, distribute, or use existing stocks of canceled products until supplies are exhausted, provided that such sale, distribution, or use is consistent with the terms of the previously approved labeling on, or that accompanied, the canceled products.

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

Dated: July 15, 2020.

Delores Barber,

Director, Information Technology and Resources Management Division, Office of Pesticide Programs.

[FR Doc. 2020–16461 Filed 7–29–20; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2020–0306; FRL–10011–30]

Petition To Revoke All Neonicotinoid Tolerances; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA seeks public comment on a May 4, 2020 petition by the Natural Resources Defense Council (NRDC) requesting that the Agency revoke all tolerances for residues of the neonicotinoid pesticides acetamiprid, clothianidin, dinotefuran, imidacloprid, and thiamethoxam. The petitioners claim that the underlying analysis supporting these tolerances are flawed and that proper consideration of available data demonstrates that the tolerances are not safe and must be revoked. A copy of the petition is available at *regulations.gov* in docket ID EPA–HQ–OPP–2020–0306.

DATES: Comments must be received on or before August 31, 2020.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2020–0306, 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:* OPP 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.*

Please note that due to the public health emergency the EPA Docket Center (EPA/DC) and Reading Room was closed to public visitors on March 31, 2020. Our EPA/DC staff will continue to provide customer service via email, phone, and webform. For further information on EPA/DC services, docket contact information and the current status of the EPA/DC and Reading Room, please visit *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Jonathan Williams, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (703)–347–0670; email address: *williams.jonathanr@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 those involved with pesticide manufacture, sale, or use; to a member or affiliate of an agricultural trade or interest group, an environmental interest group, or a public health interest group; to federal, state, or local regulatory partners; or to a member of the general public interested in the manufacture, sale, or use of pesticides (including neonicotinoids). Given the broad interest, the Agency has not attempted to identify or describe all the specific entities that may be affected by this action.

The following list of North American Industry Classification System (NAICS) codes is not intended to be exhaustive, but rather provides a guide to help readers determine whether this document applies to them. Potentially affected entities may include:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

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 the 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.*

C. How can I get copies of this document and other related information?

A copy of the NRDC's Petition memorandum, *RE: Petition to Revoke All Neonic Tolerances and Comments Regarding Dietary Exposure,* is available in the docket under docket identification (ID) number EPA–HQ–OPP–2020–0306.

II. What action is the Agency taking?

EPA seeks public comment during the next [30] days on a petition (available in docket number EPA–HQ–OPP–2020–0306) received from the NRDC requesting that the Agency revoke all tolerances for residues of the neonicotinoid pesticides acetamiprid, clothianidin, dinotefuran, imidacloprid, and thiamethoxam. The petition was submitted under section 408(d) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 346a(d). The petitioners claim that the existing tolerances are not safe and must be revoked due to several flaws in EPA's analysis of neonicotinoid toxicity and exposure when conducting its human health risk and drinking water assessments for these pesticides. The petitioners claim that EPA failed to use the most sensitive endpoint and appropriate uncertainty factors, including the full 10x children's safety factor, in not considering the potential for developmental effects in children from neonicotinoid exposure and evidence of toxic effects at low exposure levels; failed to assess the potential for cumulative toxicity from exposure to multiple neonicotinoids; failed to assess the aggregate toxicity of neonicotinoids and other chemicals resulting from interactions between neonicotinoids and chemicals used in drinking water sanitation; and failed to consider risks to highly-exposed individuals in the acute dietary risk assessment. The petitioners therefore contend that the established tolerances are not conservative enough to protect the general population, and children, from exposure to toxic amounts of neonicotinoids in food.

EPA's human health and drinking water risk assessments the neonicotinoids are contained in the dockets for each of the respective registration review cases, listed here:

- *Acetamiprid:* EPA–HQ–OPP–2012–0329.
- *Clothianidin:* EPA–HQ–OPP–2011–0865.
- *Dinotefuran:* EPA–HQ–OPP–2011–0920.
- *Imidacloprid:* EPA–HQ–OPP–2008–0844.
- *Thiamethoxam:* EPA–HQ–OPP–2011–0581.

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

Dated: July 21, 2020.

Mary Reaves,

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

[FR Doc. 2020–16454 Filed 7–29–20; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0773; FRS 16947]

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.'' 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.

DATES: Written comments and recommendations for the proposed information collection should be submitted on or before August 31, 2020.

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 Nicole Ongele, FCC, via email to *PRA@fcc.gov* and to *Nicole.Ongele@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 Nicole Ongele at (202) 418–2991. 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: 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 reduce the information collection burden for small business concerns with fewer than 25 employees.''

OMB Control Number: 3060–0773.

Title: Sections 2.803 and 2.803(c)(2), Marketing of RF Devices Prior to Equipment Authorization.

Form Number: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Businesses or other for-profit.

Number of Respondents and Responses: 10,000 respondents and 10,000 responses.

Estimated Time per Response: 0.5 hours.

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

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. 154(i), 302, 303, 303(r), and 307.