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 Code of Federal Regulations (CFR) part 2.

2. *Tips for Preparing your Comments.* When submitting comments, remember to:

- Identify the notice by docket number and other identifying information (subject heading, **Federal Register** date and page number).
- Follow directions. The agency may ask you to respond to specific questions or organize comments by referencing a CFR part or section number.
- Explain why you agree or disagree; suggest alternative and substitute language for your requested changes.
- Describe any assumptions and provide any technical information and/or data that you used.
- If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.
- Provide specific examples to illustrate your concerns and suggest alternatives.
- Explain your views as clearly as possible, avoiding the use of profanity or personal threats.
- Make sure to submit your comments by the comment period deadline identified.

II. Information About the Document

Two sections of the Clean Air Act (CAA) govern the establishment and revision of the NAAQS. Section 108 (42 U.S.C. 7408) directs the Administrator to identify and list certain air pollutants and then to issue air quality criteria for those pollutants. The Administrator is to list those air pollutants that in her ''judgment, cause or contribute to air pollution which may reasonably be anticipated to endanger public health or welfare;'' ''the presence of which in the ambient air results from numerous or diverse mobile or stationary sources;'' and ''for which * * * [the Administrator] plans to issue air quality criteria * * *.'' Air quality criteria are intended to ''accurately reflect the latest scientific knowledge useful in indicating the kind and extent of all identifiable effects on public health or welfare which may be expected from the presence of [a] pollutant in the ambient air * * * '' (42 U.S.C. 7408(b)). Under section 109 (42 U.S.C. 7409), the EPA establishes primary (health-based) and secondary (welfare-based) NAAQS for pollutants for which air quality criteria are issued. Section 109(d) requires periodic review and, if appropriate, revision of existing air quality criteria. The revised air quality criteria reflect advances in scientific knowledge on the effects of the pollutant on public health or welfare. The EPA is also required to periodically review and revise the NAAQS, if appropriate, based on the revised criteria. Section 109(d)(2) requires that an independent scientific review committee ''shall complete a review of the criteria * * * and the national primary and secondary ambient air quality standards * * * and shall recommend to the Administrator any new * * * standards and revisions of the existing criteria and standards as may be appropriate * * *.'' Since the early 1980s, this independent review function has been performed by the CASAC.

Presently, the EPA is reviewing the primary NAAQS for NO_2. The first draft *Integrated Science Assessment for Oxides of Nitrogen (Health Criteria)* (ISA) was released on November 22, 2013 (78 FR 70040), and the draft *Integrated Review Plan for the Primary NAAQS for Nitrogen Dioxide* (IRP) was released on February 6, 2014 (79 FR 7184). Both documents were reviewed by the CASAC at a public meeting in March 2014, announced in a separate notice (79 FR 8701, February 13, 2014). The final IRP was released in June 2014 (79 FR 36801, June 30, 2014) and is available at *http://www.epa.gov/ttn/naaqs/standards/nox/s_nox_2012_pd.html.* The second draft ISA was made available to both the CASAC and the public (80 FR 5110, January 30, 2015), and was reviewed in addition to the *Risk and Exposure Assessment Planning Document* (REA Planning Document) (80 FR 27304, May 13, 2015) at a public meeting in June 2015 (80 FR 22993, April 24, 2015). The final ISA was then released in January 2016 (81 FR 4910, January 28, 2016) after taking into consideration the CASAC's advice and public comments.

The PA, when final, will serve to ''bridge the gap'' between the scientific information and the judgments required of the Administrator in determining whether to retain or revise the existing primary NAAQS for NO_2, and, if revision is considered, what revisions may be appropriate. The draft PA announced today builds upon information presented in the final ISA and the REA Planning Document. The draft PA will be available on or about September 16, 2016, through the agency's Technology Transfer Network (TTN) Web site at *https://www3.epa.gov/ttn/naaqs/standards/nox/s_nox_index.html.*

The EPA is soliciting advice and recommendations from the CASAC by means of a review of this draft document at an upcoming public meeting of the CASAC, scheduled for November 9–10, 2016. Information about this public meeting will be published as a separate notice in the **Federal Register**. Following the CASAC meeting, the EPA will consider comments received from the CASAC and the public in preparing revisions to this document. The EPA will also consider public comments submitted in response to this notice when revising the document. Comments should be submitted to the docket, as described above. The document that is the subject of today's notice does not represent and should not be construed to represent any final EPA policy, viewpoint or determination.

Dated: September 15, 2016.

Stephen Page,

Director, Office of Air Quality Planning and Standards.

[FR Doc. 2016–22681 Filed 9–21–16; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2009–0317; FRL–9952–53]

Registration Review; Draft Malathion Human Health Risk Assessment; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's draft human health risk assessment for the registration review of malathion (case 0248) for public review and comment. Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. As part of the registration review process, the Agency has completed a comprehensive draft human health risk assessment for malathion. After reviewing comments received during the public comment period, EPA may issue a revised human health risk assessment, explain any changes to the draft risk assessment, respond to comments, and may request public input on risk mitigation before completing its proposed registration

review decision for malathion. Through this program, EPA is ensuring that each pesticide's registration is based on current scientific and other knowledge, including its effects on human health and the environment.

DATES: Comments must be received on or before November 21, 2016.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2009–0317, 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.* Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is available at *http://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT:

For pesticide specific information contact the Chemical Review Manager: Steven Snyderman at telephone number: (703) 347–0249; email address: *snyderman.steven@epa.gov.*

For general questions on the registration review program, contact: Richard Dumas, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 308–8015; email address: *dumas.richard@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 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 identified in **FOR FURTHER INFORMATION CONTACT**.

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 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.*

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 pesticide discussed in this document, compared to the general population.

II. Authority

EPA is conducting its registration review of malathion pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) (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 the registrations of pesticides are to be reviewed every 15 years. Under FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5) (7 U.S.C. 136a(c)(5)). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

III. Registration Review

As directed by FIFRA section 3(g), EPA is reviewing the pesticide registrations for malathion to ensure that they continue to satisfy the FIFRA standard for registration—that is, that malathion can still be used without unreasonable adverse effects on human health or the environment. Information concerning the registration review of malathion (case 0248) is in the docket, under Docket ID No. EPA–HQ–OPP–2009–0317.

Pursuant to 40 CFR 155.53(c), EPA is providing an opportunity, through this notice of availability, for interested parties to provide comments and input concerning the Agency's draft human health risk assessment for malathion. Such comments and input could address, among other things, the Agency's risk assessment methodologies and assumptions, as applied to this draft human health risk assessment. The Agency will consider all comments received during the public comment period and make changes, as appropriate, to the draft human health risk assessment. EPA will then issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments. In the **Federal Register** notice announcing the availability of the revised risk assessment, if the revised risk assessment indicates risks of concern, the Agency may provide a comment period for the public to submit suggestions for mitigating the risk identified in the revised risk assessment before developing a proposed registration review decision for malathion.

1. *Other related information.* Additional information on the registration review status of malathion, as well as information on the Agency's registration review program and on its implementing regulation is available at *https://www.epa.gov/pesticide-reevaluation.*

2. *Information submission requirements.* Anyone may submit data or information in response to this document. To be considered during a pesticide's registration review, the submitted data or information must meet the following requirements:

- To ensure that EPA will consider data or information submitted, interested persons must submit the data or information during the comment

period. The Agency may, at its discretion, consider data or information submitted at a later date.

- The data or information submitted must be presented in a legible and useable form. For example, an English translation must accompany any material that is not in English and a written transcript must accompany any information submitted as an audiographic or videographic record. Written material may be submitted in paper or electronic form.
- Submitters must clearly identify the source of any submitted data or information.
- Submitters may request the Agency to reconsider data or information that the Agency rejected in a previous review. However, submitters must explain why they believe the Agency should reconsider the data or information in the pesticide's registration review.

As provided in 40 CFR 155.58, the registration review docket for each pesticide case will remain publicly accessible through the duration of the registration review process; that is, until all actions required in the final decision on the registration review case have been completed.

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

Dated: September 14, 2016.

Yu-Ting Guilaran,

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

[FR Doc. 2016–22881 Filed 9–21–16; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0653]

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 (44 U.S.C. 3501–3520), 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.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid 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 PRA comments should be submitted on or before November 21, 2016. 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 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:

OMB Control Number: 3060–0653.

Title: Sections 64.703(b) and (c), Consumer Information—Posting by Aggregators.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for-profit entities.

Number of Respondents: 56,075 respondents; 5,339,038 responses.

Estimated Time per Response: .017 hours (1 minute) to 3 hours.

Frequency of Response: On occasion reporting requirements; Third party disclosure.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this information collection is found at section 226 [47 U.S.C. 226] Telephone Operator Services codified at 47 CFR 64.703(b) Consumer Information.

Total Annual Burden: 174,401 hours.

Total Annual Cost: $1,343,721.

Privacy Act Impact Assessment: An assurance of confidentiality is not offered because this information collection does not require the collection of personally identifiable information (PII) from individuals.

Nature and Extent of Confidentiality: No impact(s).

Needs and Uses: The information collection requirements included under this OMB Control Number 3060–0653, requires aggregators (providers of telephones to the public or to transient users of their premises) under 47 U.S.C. 226(c)(1)(A), 47 CFR 64.703(b) of the Commission's rules, to post in writing, on or near such phones, information about the pre-subscribed operator services, rates, carrier access, and the FCC address to which consumers may direct complaints.

Section 64.703(c) of the Commission's rules requires the posted consumer information to be added when an aggregator has changed the pre-subscribed operator service provider (OSP) no later than 30 days following such change. Consumers will use this information to determine whether they wish to use the services of the identified OSP.

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2016–22797 Filed 9–21–16; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0655]

Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority

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 (44 U.S.C. 3501–3520), 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,