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III. How to Submit Technical Comments to the Docket

Please note that questions or comments regarding the peer review process, including comments on the candidate pool of peer reviewers, should be directed to chloroprenePBPk@versar.com (Subject line: Chloroprene PBPk Peer Review); or by phone (301) 304-3121 (ask for Tracey Cowen). Comments on the technical documents related to the PBPk modeling should be submitted to Docket ID No. EPA-HQ-ORD-2020-0181, by one of the following methods:

- **Online:** <https://www.regulations.gov>. Follow the online instructions for submitting comments.
- **Email:** Docket_ORD@epa.gov.
- **Fax:** 202-566-9744. Due to COVID-19, there may be a delay in processing comments submitted by fax.
- **Mail:** U.S. Environmental Protection Agency, EPA Docket Center (ORD Docket), Mail Code: 28221T, 1200 Pennsylvania Avenue NW, Washington, DC 20460. The phone number is 202-566-1752. Due to COVID-19, there may be a delay in processing comments submitted by mail.

Note: The EPA Docket Center and Reading Room is currently closed to public visitors to reduce the risk of transmitting COVID-19. Docket Center staff will continue to provide remote customer service via email, phone, and webform. The public can submit comments via www.Regulations.gov or email. No hand deliveries are currently being accepted.

Instructions: Direct your comments to Docket ID No. EPA-HQ-ORD-2020-0181. Please ensure that your comments are submitted within the specified comment period. Comments received after the closing date will be marked "late," and may only be considered if time permits. It is EPA's policy to include all comments it receives in the public docket without change and to make the comments available online at www.regulations.gov, including any personal information provided, unless a comment includes information claimed to be Confidential Business Information (CBI) or other information for which disclosure is restricted by statute. Do not submit information through www.regulations.gov or email that you consider to be CBI or otherwise protected. The www.regulations.gov website is an "anonymous access"

system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an email comment directly to EPA without going through www.regulations.gov, your email address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD-ROM you submit. 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. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses. For additional information about EPA's public docket visit the EPA Docket Center homepage at www.epa.gov/epahome/dockets.htm.

Docket: Documents in the docket are listed in the www.regulations.gov index. Although listed in the index, some information is not publicly available, e.g., CBI or other information whose disclosure is restricted by statute. Certain other materials, such as copyrighted material, are publicly available only in hard copy. Publicly available docket materials are available either electronically in www.regulations.gov or in hard copy at the ORD Docket in the EPA Headquarters Docket Center.

Wayne E. Cascio,
Director, Center for Public Health & Environmental Assessment.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-ORD-2020-0183; FRL-10012-20-ORD]

Availability of the IRIS Assessment Plan for Oral Exposure to Vanadium and Compounds

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of public comment period.

SUMMARY: The Environmental Protection Agency (EPA) is announcing a 30-day public comment period associated with release of the IRIS Assessment Plan for Oral Exposure to Vanadium and Compounds. This document communicates information on the

scoping needs identified by EPA program and regional offices and the IRIS Program's initial problem formulation activities. Specifically, the assessment plan outlines the objectives for the IRIS assessment and the type of evidence considered most pertinent to address the scoping needs. EPA is releasing this IRIS Assessment Plan for a 30-day public comment period in advance of a public science webinar planned for August 19, 2020. The Agency encourages the public to comment on all aspects of the assessment plan, including key science issues and identification of any new or missing studies.

DATES: The 30-day public comment period begins July 24, 2020 and ends August 24, 2020. Comments must be received on or before August 24, 2020.

ADDRESSES: The IRIS Assessment Plan for Oral Exposure to Vanadium and Compounds will be available via the internet on the IRIS website at <https://www.epa.gov/iris/iris-recent-additions> and in the public docket at <http://www.regulations.gov>, Docket ID No. EPA-HQ-ORD-2020-0182.

FOR FURTHER INFORMATION CONTACT: For information on the public comment period, contact the ORD Docket at the EPA Headquarters Docket Center; **telephone:** 202-566-1752; **facsimile:** 202-566-9744; or **email:** Docket_ORD@epa.gov.

For technical information on the IRIS Assessment Plan for Oral Exposure to Vanadium and Compounds, contact Dr. James Avery, CPHEA; **telephone:** 202-564-1494; or **email:** avery.james@epa.gov.

SUPPLEMENTARY INFORMATION:

I. Information About IRIS Assessment Plans

EPA's IRIS Program is a human health assessment program that evaluates quantitative and qualitative information on the health effects that may result from exposure to chemicals found in the environment. Through the IRIS Program, EPA provides high quality science-based human health assessments to support the Agency's regulatory activities and decisions to protect public health. As part of scoping and initial problem formulation activities prior to the development of an assessment, the IRIS Program carries out a broad, preliminary literature survey to assist in identifying health effects that have been studied in relation to the chemical or substance of interest, as well as science issues that may need to be considered when evaluating toxicity. This information, in conjunction with scoping needs identified by EPA

program and regional offices, is used to inform the development of an IRIS Assessment Plan (IAP).

The IAP communicates the plan for developing each individual chemical assessment to the public and includes summary information on the IRIS Program's scoping and initial problem formulation activities, objectives and specific aims for the assessment, and a PECO (Populations, Exposures, Comparators, and Outcomes) for the systematic review. The PECO provides the framework for developing detailed literature search strategies and inclusion/exclusion criteria, particularly with respect to evidence stream (e.g., human, animal, mechanistic), exposure measures, and outcome measures. The IAP serves to inform the subsequent development of chemical-specific systematic review protocols, which will be made available for public review.

II. Public Webinar Information

To allow for public input, EPA is convening a public webinar to discuss the IRIS Assessment Plan for Oral Exposure to Vanadium and Compounds on August 19, 2020. Specific teleconference and webinar information regarding this public meeting will be provided through the IRIS website (<https://www.epa.gov/iris>) and via EPA's IRIS listserv. To register for the IRIS listserv, visit the IRIS website (<https://www.epa.gov/iris>) or visit <https://www.epa.gov/iris/forms/staying-connected-integrated-risk-information-system#connect>.

III. How to Submit Technical Comments to the Docket at <https://www.regulations.gov>

Submit your comments, identified by Docket ID No. EPA-HQ-ORD-2020-0183 for IRIS Assessment Plan for Oral Exposure to Vanadium and Compounds, by one of the following methods:

- www.regulations.gov: Follow the on-line instructions for submitting comments.
- *Email*: Docket_ORD@epa.gov.
- *Fax*: 202-566-9744. Due to COVID-19, there may be a delay in processing comments submitted by fax.
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customer service via email, phone, and webform. The public can submit comments via www.Regulations.gov or email. No hand deliveries are currently being accepted.

Instructions: Direct your comments to Docket ID No. EPA-HQ-ORD-2020-0183. Please ensure that your comments are submitted within the specified comment period. Comments received after the closing date will be marked "late," and may only be considered if time permits. It is EPA's policy to include all comments it receives in the public docket without change and to make the comments available online at www.regulations.gov, including any personal information provided, unless a comment includes information claimed to be Confidential Business Information (CBI) or other information for which disclosure is restricted by statute. Do not submit information through www.regulations.gov or email that you consider to be CBI or otherwise protected. The www.regulations.gov website is an "anonymous access" system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an email comment directly to EPA without going through www.regulations.gov, your email address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD-ROM you submit. 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. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses. For additional information about EPA's public docket visit the EPA Docket Center homepage at www.epa.gov/epahome/dockets.htm.

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the ORD Docket in the EPA Headquarters Docket Center.

Wayne E. Cascio,

Director, Center for Public Health & Environmental Assessment.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OGC-2020-0368; FRL-10012-76-OGC]

Proposed Consent Decree, Clean Air Act Citizen Suit

AGENCY: Environmental Protection Agency (EPA)

ACTION: Notice of proposed consent decree; request for public comment.

SUMMARY: In accordance with section 113(g) of the Clean Air Act, as amended ("CAA" or the "Act"), notice is given of a proposed consent decree in *Our Children's Earth Foundation v. Wheeler*, No. 3:19-cv-07125 (N.D. Cal.). On October 29, 2019, Our Children's Earth Foundation filed a complaint in the United States District Court for the Northern District of California, alleging that the Administrator of the United States Environmental Protection Agency ("EPA") failed to perform non-discretionary duties to review the existing New Source Performance Standards ("NSPS") governing Secondary Lead Smelters ("Secondary Lead Smelters NSPS"); Lead-Acid Battery Manufacturing Plants ("Lead-Acid Battery Manufacturing NSPS"); Industrial Surface Coating: Surface Coating of Plastic Parts for Business Machines ("Industrial Surface Coating of Plastic Parts for Business Machines NSPS"); and Automobile and Light Duty Truck Surface Coating Operations ("Automobile and Light Duty Truck Surface Coating Operations NSPS"), and to review the existing National Emission Standards for Hazardous Air Pollutants ("NESHAP") governing Dry Cleaning Facilities: National Perchloroethylene Air Emission Standards ("Dry Cleaning Facilities NESHAP"); Paint Stripping and Miscellaneous Surface Coating Operations at Area Sources ("Paint Stripping and Miscellaneous Surface Coating Operations NESHAP"), and Lead Acid Battery Manufacturing Area Sources ("Lead Acid Battery Manufacturing NESHAP"). The proposed consent decree would establish deadlines for EPA to take action on these source categories.