

eRegistration link. Select the eFiling link to log on and submit the intervention or protests.

Persons unable to file electronically may mail similar pleadings to the Federal Energy Regulatory Commission, 888 First Street NE, Washington, DC 20426. Hand delivered submissions in docketed proceedings should be delivered to Health and Human Services, 12225 Wilkins Avenue, Rockville, Maryland 20852.

In addition to publishing the full text of this document in the **Federal Register**, the Commission provides all interested persons an opportunity to view and/or print the contents of this document via the internet through the Commission's Home Page (<http://ferc.gov>) using the "eLibrary" link. Enter the docket number excluding the last three digits in the docket number field to access the document. At this time, the Commission has suspended access to the Commission's Public Reference Room, due to the proclamation declaring a National Emergency concerning the Novel Coronavirus Disease (COVID-19), issued by the President on March 13, 2020. For assistance, contact the Federal Energy Regulatory Commission at [FERCOnlineSupport@ferc.gov](mailto:FERCOnlineSupport@ferc.gov) or call toll-free, (886) 208-3676 or TTY, (202) 502-8659.

Dated: December 14, 2020.

**Kimberly D. Bose,**

*Secretary.*

[FR Doc. 2020-27932 Filed 12-17-20; 8:45 am]

**BILLING CODE 6717-01-P**

## ENVIRONMENTAL PROTECTION AGENCY

[ER-FRL-9054-4]

### Environmental Impact Statements; Notice of Availability

*Responsible Agency:* Office of Federal Activities, General Information 202-564-5632 or <https://www.epa.gov/nepa>. Weekly receipt of Environmental Impact Statements (EIS)

Filed December 7, 2020 10 a.m. EST  
Through December 14, 2020 10 a.m. EST

Pursuant to 40 CFR 1506.9.

*Notice:* Section 309(a) of the Clean Air Act requires that EPA make public its comments on EISs issued by other Federal agencies. EPA's comment letters on EISs are available at: <https://cdxnodengn.epa.gov/cdx-enepa-public/action/eis/search>.

EIS No. 20200260, Final Supplement, BR, CA, B.F. Sisk Dam Raise and

Reservoir Expansion Project, Review Period Ends: 01/19/2021, Contact: Casandra Arthur 530-892-6202.

EIS No. 20200261, Draft Supplement, USACE, FL, Port Everglades Harbor, Broward County, Florida, Comment Period Ends: 02/01/2021, Contact: Angela Dunn 904-232-2336.

### Amended Notice

EIS No. 20200210, Draft, STB, UT, Uinta Basin Railway, Comment Period Ends: 01/28/2021, Contact: Joshua Wayland 202-245-0330.

Revision to FR Notice Published 10/30/2020; Extending the Comment Period from 12/14/2020 to 01/28/2021.

EIS No. 20200242, Draft, USACE, VA, Surry To Skiffes Creek to Whealton Transmission Project, Comment Period Ends: 02/10/2021, Contact: Randy Steffey 757-201-7579.

Revision to FR Notice Published 11/27/2020; Extending the Comment Period from 01/11/2021 to 02/10/2021.

Dated: December 14, 2020.

**Cindy S. Barger,**

*Director, NEPA Compliance Division, Office of Federal Activities.*

[FR Doc. 2020-27888 Filed 12-17-20; 8:45 am]

**BILLING CODE 6560-50-P**

## ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPPT-2019-0502; FRL-10017-44]

### Perchloroethylene (PCE); Final Toxic Substances Control Act (TSCA) Risk Evaluation; Notice of Availability

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** The Environmental Protection Agency (EPA) is announcing the availability of the final Toxic Substances Control Act (TSCA) risk evaluation of perchloroethylene (PCE). The purpose of conducting risk evaluations under TSCA is to determine whether a chemical substance presents an unreasonable risk of injury to health or the environment under the conditions of use, including an unreasonable risk to a relevant potentially exposed or susceptible subpopulation, without consideration of costs or other nonrisk factors. EPA has determined that specific conditions of use of PCE present an unreasonable risk of injury to health or the environment. For those conditions of use for which EPA has found an unreasonable risk, EPA must move to address that unreasonable risk through risk management measures enumerated in

TSCA. EPA has also determined that specific conditions of use do not present unreasonable risk of injury to health or the environment. For those conditions of use for which EPA has found no unreasonable risk to health or the environment, the Agency's determination is a final Agency action and is issued via order in the risk evaluation.

**ADDRESSES:** The docket for this action, identified by docket identification (ID) number EPA-HQ-OPPT-2019-0502, is available online at <http://www.regulations.gov> or in-person at the Office of Pollution Prevention and Toxics Docket (OPPT Docket), Environmental Protection Agency Docket Center (EPA/DC), West William Jefferson Clinton Bldg., Rm. 3334, 1301 Constitution Ave. NW, Washington, DC. The 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, and the telephone number for the OPPT Docket is (202) 566-0280.

Due to the public health concerns related to COVID-19, the EPA Docket Center (EPA/DC) and Reading Room is closed to visitors with limited exceptions. The staff continues to provide remote customer service via email, phone, and webform. For the latest status information on EPA/DC services and docket access, visit <https://www.epa.gov/dockets>.

**FOR FURTHER INFORMATION CONTACT:** For technical information contact: Yvette Selby-Mohamadu, Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460-0001; telephone number: (202) 564-5245; email address: [selby-mohamadu.yvette@epa.gov](mailto:selby-mohamadu.yvette@epa.gov).

For general information contact: The TSCA-Hotline, ABVI-Goodwill, 422 South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554-1404; email address: [TSCA-Hotline@epa.gov](mailto:TSCA-Hotline@epa.gov).

### SUPPLEMENTARY INFORMATION:

#### I. General Information

##### A. Does this action apply to me?

This action is directed to the public in general. This action may be of interest to persons who are or may be interested in risk evaluations of chemical substances under TSCA, 15 U.S.C. 2601 *et seq.* Since other entities may also be interested in this final risk evaluation, the EPA has not attempted to describe all the specific entities that may be affected by this action.

*B. What is the Agency's authority for taking this action?*

TSCA section 6, 15 U.S.C. 2605, requires EPA to conduct risk evaluations to “determine whether a chemical substance presents an unreasonable risk of injury to health or the environment, without consideration of costs or other nonrisk factors, including an unreasonable risk to a potentially exposed or susceptible subpopulation identified as relevant to the risk evaluation by the Administrator, under the conditions of use.” 15 U.S.C. 2605(b)(4)(A). TSCA sections 6(b)(4)(A) through (H) enumerate the deadlines and minimum requirements applicable to this process, including provisions that provide instruction on chemical substances that must undergo evaluation, the minimum components of a TSCA risk evaluation, and the timelines for public comment and completion of the risk evaluation. TSCA also requires that EPA operate in a manner that is consistent with the best available science, make decisions based on the weight of the scientific evidence and consider reasonably available information. 15 U.S.C. 2625(h), (i), and (k). TSCA section 6(i) directs that a determination of “no unreasonable risk” shall be issued by order and considered to be a final Agency action, while a determination of “unreasonable risk” is not considered to be a final Agency action. 15 U.S.C. 2605(i).

The statute identifies the minimum components for all chemical substance risk evaluations. For each risk evaluation, EPA must publish a document that outlines the scope of the risk evaluation to be conducted, which includes the hazards, exposures, conditions of use, and the potentially exposed or susceptible subpopulations that EPA expects to consider. 15 U.S.C. 2605(b)(4)(D). The statute further provides that each risk evaluation must also: (1) integrate and assess available information on hazards and exposures for the conditions of use of the chemical substance, including information that is relevant to specific risks of injury to health or the environment and information on relevant potentially exposed or susceptible subpopulations; (2) describe whether aggregate or sentinel exposures were considered and the basis for that consideration; (3) take into account, where relevant, the likely duration, intensity, frequency, and number of exposures under the conditions of use; and (4) describe the weight of the scientific evidence for the identified hazards and exposures. 15 U.S.C. 2605(b)(4)(F)(i) through (ii) and (iv) through (v). Each risk evaluation

must not consider costs or other nonrisk factors. 15 U.S.C. 2605(b)(4)(F)(iii).

The statute requires that the risk evaluation process be completed within a specified timeframe and provide an opportunity for public comment on a draft risk evaluation prior to publishing a final risk evaluation. 15 U.S.C. 2605(b)(4).

Subsection 5.4.1 of the final risk evaluation for PCE constitutes the order required under TSCA section 6(i)(1), and the “no unreasonable risk” determinations in that subsection are considered to be a final Agency action effective on the date of issuance of the order. In conducting risk evaluations, “EPA will determine whether the chemical substance presents an unreasonable risk of injury to health or the environment under each condition of use within the scope of the risk evaluation. . . .” 40 CFR 702.47. Under EPA’s implementing regulations, “[a] determination by EPA that the chemical substance, under one or more of the conditions of use within the scope of the risk evaluation, does not present an unreasonable risk of injury to health or the environment will be issued by order and considered to be a final Agency action, effective on the date of issuance of the order.” 40 CFR 702.49(d). For purposes of TSCA section 19(a)(1)(A), the date of issuance of the TSCA section 6(i)(1) order for PCE shall be at 1:00 p.m. Eastern time (standard or daylight, as appropriate) on the date that is two weeks after the date when this notice is published in the **Federal Register**, which is in accordance with 40 CFR 23.5.

*C. What action is EPA taking?*

EPA is announcing the availability of the risk evaluation of the chemical substance identified in Unit II. In this risk evaluation EPA has made unreasonable risk determinations on some of the conditions of use within the scope of the risk evaluation for this chemical. For those conditions of use for which EPA has found an unreasonable risk of injury to health or the environment, EPA must initiate regulatory action to address those risks through risk management measures enumerated in 15 U.S.C. 2605(a).

EPA also is announcing the availability of the information required to be provided publicly with each risk evaluation, which is available online at <http://www.regulations.gov> in the dockets identified. 40 CFR 702.51. Specifically, EPA has provided:

- The scope document and problem formulation (in Docket ID No. EPA–HQ–OPPT–2016–0732);

- Draft risk evaluation and final risk evaluation (in Docket ID No. EPA–HQ–OPPT–2019–0502);

- All notices, determinations, findings, consent agreements, and orders (in Docket ID No. EPA–HQ–OPPT–2019–0502);

- Any information required to be provided to the Agency under 15 U.S.C. 2603 (in Docket ID No. EPA–HQ–OPPT–2016–0732 and Docket ID No. EPA–HQ–OPPT–2019–0502);

- A nontechnical summary of the risk evaluation (in Docket ID No. EPA–HQ–OPPT–2019–0502);

- A list of the studies, with the results of the studies, considered in carrying out each risk evaluation (Risk Evaluation for Perchloroethylene (Ethene, 1,1,2,2-Tetrachloro-) in Docket ID No. EPA–HQ–OPPT–2019–0502);

- The final peer review report, including the response to peer review and public comments received during peer review (in Docket ID No. EPA–HQ–OPPT–2019–0502); and

- Response to public comments received on the draft scope and the draft risk evaluation (in Docket ID No. EPA–HQ–OPPT–2019–0502).

## II. TSCA Risk Evaluation

*A. What is EPA's risk evaluation process for existing chemicals under TSCA?*

The risk evaluation process is the second step in EPA’s existing chemical process under TSCA, following prioritization and before risk management. As this chemical is one of the first ten chemical substances undergoing risk evaluation, the chemical substance was not required to go through prioritization (81 FR 91927, December 19, 2016) (FRL–9956–47). The purpose of conducting risk evaluations is to determine whether a chemical substance presents an unreasonable risk of injury to health or the environment under the conditions of use, including an unreasonable risk to a relevant potentially exposed or susceptible subpopulation. As part of this process, EPA must evaluate both hazard and exposure, not consider costs or other nonrisk factors, use reasonably available information and approaches in a manner that is consistent with the requirements in TSCA for the use of the best available science, and ensure decisions are based on the weight of the scientific evidence.

The specific risk evaluation process that EPA has established by rule to implement the statutory process is set out in 40 CFR part 702 and summarized on EPA’s website at <http://www.epa.gov/assessing-and-managing-chemicals-under-tsca/risk-evaluations->

*existing-chemicals-under-tsca*. As explained in the preamble to EPA's final rule on procedures for risk evaluation (82 FR 33726, July 20, 2017) (FRL-9964-38), the specific regulatory process set out in 40 CFR part 702, subpart B will be followed for the first ten chemical substances undergoing risk evaluation to the maximum extent practicable.

Prior to the publication of this final risk evaluation, a draft risk evaluation was subject to peer review and public comment. EPA reviewed the peer review report from the Science Advisory Committee on Chemicals (SACC) and public comments and has supplemented the risk evaluation in response to these comments as appropriate. The public comments and peer review report are in Docket EPA-HQ-OPPT-2019-0502 at [www.regulations.gov](http://www.regulations.gov). Prior to the publication of the draft risk evaluation, EPA made available the scope and problem formulation, and solicited public input on uses and exposure. EPA's documents and the public comments are in Docket EPA-HQ-OPPT-2019-0502. Additionally, information about the scope, problem formulation, and draft risk evaluation phases of the TSCA risk evaluation for this chemical is available at <https://www.epa.gov/assessing-and-managing-chemicals-under-tsca/risk-evaluation-perchloroethylene>.

#### B. What is Perchloroethylene?

Perchloroethylene is currently manufactured, processed, distributed, used, and disposed of as part of a wide range of industrial, commercial, and consumer conditions of use, including production of fluorinated compounds, and as a solvent in dry cleaning and vapor degreasing. Consumer and commercial products that contain perchloroethylene include adhesives (arts and crafts, as well as light repairs), aerosol degreasing, brake cleaners, aerosol lubricants, sealants, stone polish, stainless steel polish and other cleaners used for wiping surfaces. The yearly aggregate production volume for perchloroethylene ranged from 388 to 324 million pounds between 2012 and 2015 according to CDR data.

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

**Andrew Wheeler,**  
Administrator.

[FR Doc. 2020-27880 Filed 12-17-20; 8:45 am]

**BILLING CODE 6560-50-P**

## FEDERAL COMMUNICATIONS COMMISSION

[FRS 17305]

### Federal Advisory Committee Act; Technological Advisory Council

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice of public meeting.

**SUMMARY:** In accordance with the Federal Advisory Committee Act, this notice advises interested persons that the Federal Communications Commission's (FCC) Technological Advisory Council will hold a meeting on Thursday January 14, 2021 via conference call and available to the public via the internet at <http://www.fcc.gov/live>, from 10:00 a.m. to 12:30 p.m.

**DATES:** Thursday January 14, 2021.

**ADDRESSES:** Federal Communications Commission, 45 L Street NE, Washington, DC 20554.

**FOR FURTHER INFORMATION CONTACT:** Michael Ha, Deputy Chief, Policy and Rules Division 202-418-2099; [michael.ha@fcc.gov](mailto:michael.ha@fcc.gov).

**SUPPLEMENTARY INFORMATION:** At the January 14th meeting, the TAC will consider and vote on a white paper prepared by the Artificial Intelligence working group and recommendations from its four working groups: 5G/IOT/V-RAN, Future of Unlicensed Operations, Artificial Intelligence, and 5G Radio Access Network Technology. This agenda may be modified at the discretion of the TAC Chair and the Designated Federal Officer (DFO). Meetings are broadcast live with open captioning over the internet from the FCC Live web page at <http://www.fcc.gov/live/>. The public may submit written comments before the meeting to Michael Ha, the FCC's Designated Federal Officer for Technological Advisory Council by email: [michael.ha@fcc.gov](mailto:michael.ha@fcc.gov) or U.S. Postal Service Mail (Michael Ha, Federal Communications Commission, 45 L Street NE, Washington, DC 20554). Open captioning will be provided for this event. Other reasonable accommodations for people with disabilities are available upon request. Requests for such accommodations should be submitted via email to [fcc504@fcc.gov](mailto:fcc504@fcc.gov) or by calling the Office of Engineering and Technology at 202-418-2470 (voice), (202) 418-1944 (fax). Such requests should include a detailed description of the accommodation needed. In addition, please include your contact information. Please allow at least five days advance notice; last

minute requests will be accepted but may not be possible to fill.

Federal Communications Commission.

**Ronald T. Repasi,**

*Acting Chief, Office of Engineering and Technology.*

[FR Doc. 2020-27832 Filed 12-17-20; 8:45 am]

**BILLING CODE 6712-01-P**

## FEDERAL FINANCIAL INSTITUTIONS EXAMINATION COUNCIL

[Docket No. AS20-14]

### Agency Information Collection Activities; Renewal of an Approved Information Collection: Reporting information for the AMC Registry

**AGENCY:** Appraisal Subcommittee of the Federal Financial Institutions Examination Council (ASC)

**ACTION:** Notice and request for comments.

**SUMMARY:** In accordance with the Paperwork Reduction Act of 1995, the ASC invites public comments on our intention to request the Office of Management and Budget (OMB) approval to renew an information collection request entitled "Reporting information for the AMC Registry."

**DATES:** Written comments must be received on or before February 16, 2021 to be assured of consideration.

**ADDRESSES:** Commenters are encouraged to submit comments by the Federal eRulemaking Portal or email, if possible. You may submit comments, identified by Docket Number AS20-14, by any of the following methods:

- **Federal eRulemaking Portal:** <https://www.Regulations.gov>. Follow the instructions for submitting comments. Click on the "Help" tab on the *Regulations.gov* home page to get information on using *Regulations.gov*, including instructions for submitting public comments.
- **E-Mail:** [webmaster@asc.gov](mailto:webmaster@asc.gov). Include the docket number in the subject line of the message.
- **Fax:** (202) 289-4101. Include the docket number of fax cover sheet.
- **Mail:** Address to Appraisal Subcommittee, Attn: Lori Schuster, Management and Program Analyst, 1325 G Street NW, Suite 500, Washington, DC 20005.
- **Hand Delivery/Courier:** 1325 G Street NW, Suite 500, Washington, DC 20005.

In general, the ASC will enter all comments received into the docket and publish those comments on the *Regulations.gov* website without change, including any business or