proclamation declaring a National Emergency concerning the Novel Coronavirus Disease (COVID–19), issued by the President on March 13, 2020. For assistance, contact FERC Online Support.

You may also register online at *http://www.ferc.gov/docs-filing/esubscription.asp* to be notified via email of new filings and issuances related to this or other pending projects. For assistance, contact FERC Online Support.

Dated: April 7, 2023.

Kimberly D. Bose,

Secretary.

[FR Doc. 2023–07806 Filed 4–12–23; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission

Combined Notice of Filings

Take notice that the Commission has received the following Natural Gas & Oil Pipeline Rate and Refund Report filings:

Filings in Existing Proceedings

Docket Numbers: RP23–621–000.

Applicants: Northern Border Pipeline Company.

Description: Report Filing: Supplement to NBPL 2023 CUS Filing to be effective N/A.

Filed Date: 4/6/23.

Accession Number: 20230406–5139.

Comment Date: 5 p.m. ET 4/18/23.

Any person desiring to protest in any the above proceedings must file in accordance with Rule 211 of the Commission's Regulations (18 CFR 385.211) on or before 5:00 p.m. Eastern time on the specified comment date.

The filings are accessible in the Commission's eLibrary system (*https://elibrary.ferc.gov/idmws/search/fercgensearch.asp*) by querying the docket number.

eFiling is encouraged. More detailed information relating to filing requirements, interventions, protests, service, and qualifying facilities filings can be found at: *http://www.ferc.gov/docs-filing/efiling/filing-req.pdf.* For other information, call (866) 208–3676 (toll free). For TTY, call (202) 502–8659.

Dated: April 7, 2023.

Debbie-Anne A. Reese,

Deputy Secretary.

[FR Doc. 2023–07777 Filed 4–12–23; 8:45 am]

BILLING CODE 6717–01–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–10855–01–OMS]

Cross-Media Electronic Reporting: Authorized Program Revision Approval, New Jersey Department of Environmental Protection (NJDEP)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the Environmental Protection Agency's (EPA) approval of the New Jersey Department of Environmental Protection (NJDEP) request to revise/modify certain of its EPA-authorized programs to allow electronic reporting.

DATES: EPA approves the authorized program revisions/modifications as of April 13, 2023.

FOR FURTHER INFORMATION CONTACT: Shirley M. Miller, U.S. Environmental Protection Agency, Office of Information Management, Mail Stop 2824T, 1200 Pennsylvania Avenue NW, Washington, DC 20460, (202) 566–2908, *miller.shirley@epa.gov.*

SUPPLEMENTARY INFORMATION: On October 13, 2005, the final Cross-Media Electronic Reporting Rule (CROMERR) was published in the **Federal Register** (70 FR 59848) and codified as part 3 of title 40 of the CFR. CROMERR establishes electronic reporting as an acceptable regulatory alternative to paper reporting and establishes requirements to assure that electronic documents are as legally dependable as their paper counterparts. Subpart D of CROMERR requires that state, tribal or local government agencies that receive, or wish to begin receiving, electronic reports under their EPA-authorized programs must apply to EPA for a revision or modification of those programs and obtain EPA approval. Subpart D provides standards for such approvals based on consideration of the electronic document receiving systems that the state, tribe, or local government will use to implement the electronic reporting. Additionally, § 3.1000(b) through (e) of 40 CFR part 3, subpart D provides special procedures for program revisions and modifications to allow electronic reporting, to be used at the option of the state, tribe or local government in place of procedures available under existing program-specific authorization regulations. An application submitted under the subpart D procedures must show that the state, tribe or local government has sufficient legal authority to implement the electronic reporting components of the programs covered by the application and will use electronic document receiving systems that meet the applicable subpart D requirements.

On March 13, 2023, the New Jersey Department of Environmental Protection (NJDEP) submitted an application titled National Pollutant Discharge Elimination System (NPDES) Electronic Reporting Tool (NeT) for revisions/modifications to its EPA-approved programs under title 40 CFR to allow new electronic reporting. EPA reviewed NJDEP's request to revise/modify its EPA-authorized programs and, based on this review, EPA determined that the application met the standards for approval of authorized program revisions/modifications set out in 40 CFR part 3, subpart D. In accordance with 40 CFR 3.1000(d), this notice of EPA's decision to approve NJDEP's request to revise/modify its following EPA-authorized programs to allow electronic reporting under 40 CFR is being published in the **Federal Register**:

Part 123: EPA-Administered Permit Programs: the National Pollutant Discharge Elimination System (NPDES) Reporting under 40 CFR 122 and 125.

NJDEP was notified of EPA's determination to approve its application with respect to the authorized programs listed above.

Dated: April 6, 2023.

Jennifer Campbell,

Director, Office of Information Management.

[FR Doc. 2023–07725 Filed 4–12–23; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2013–0244; FRL–10818–01–OCSPP]

Pesticide Registration Review; Proposed Interim Decision and Draft Risk Assessment Addendum for Ethylene Oxide; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of and solicits public comment on EPA's proposed interim registration review decision and draft risk assessment addendum for ethylene oxide.

DATES: Comments must be received on or before June 12, 2023.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2013–0244, through the *Federal eRulemaking Portal* 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 and visiting the docket, along with more information about dockets generally, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: The Chemical Review Manager for ethylene oxide as listed in table 1.

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 and human health advocates; distributors and users of medical devices; owners and operators of commercial sterilization facilities; 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 for ethylene oxide identified in table 1 in unit IV.

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 on 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: *https://www.epa.gov/dockets/commenting-epa-dockets.*

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 ethylene oxide (EtO) discussed in this document, compared to the general population.

II. Background

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 proposed interim decision and draft risk assessment addendum for ethylene oxide (table 1). 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.

III. Authority

EPA is conducting its registration review of ethylene oxide pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. Section 3(g) of FIFRA 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.

IV. What action is the Agency taking?

Pursuant to 40 CFR 155.58, this notice announces the availability of EPA's proposed interim registration review decisions for ethylene oxide and the draft risk assessment addendum and opens a 60-day public comment period on these documents.

TABLE 1—ETHYLENE OXIDE REGISTRATION REVIEW DOCKET DETAILS

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Ethylene oxide Case Number 2275	EPA–HQ–OPP–2013–0244	Jessica Bailey *OPPethyleneoxideinquiries@epa.gov.*

The registration review docket for a pesticide includes earlier documents related to the registration review case. For example, the review opened with a Preliminary Work Plan, for public comment. A Final Work Plan was placed in the docket following public comment on the Preliminary Work Plan.

The documents in the docket describe EPA's rationales for conducting additional risk assessments for the registration review of ethylene oxide, as well as the Agency's subsequent risk findings and consideration of possible risk mitigation measures. The proposed interim registration review decision is supported by the rationale included in those documents. Following public comment, the Agency will issue an interim or final registration review decision for ethylene oxide.

The registration review final rule at 40 CFR 155.58(a) provides for a minimum 60-day public comment period on all proposed interim registration review decisions. This comment period is intended to provide an opportunity for public input and a mechanism for initiating any necessary amendments to the proposed interim decision. All comments should be submitted using the methods in **ADDRESSES** and must be received by EPA on or before the closing date. These comments will become part of the docket for ethylene oxide. Comments received after the close of the comment period will be marked ''late.'' EPA is not required to consider these late comments.

The Agency will carefully consider all comments received by the closing date and may provide a ''Response to Comments Memorandum'' in the docket. The interim registration review decision will explain the effect that any comments had on the interim decision and provide the Agency's response to significant comments.

Background on the registration review program is provided at: *https://www.epa.gov/pesticide-reevaluation.*

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

Dated: March 28, 2023.

Anita Pease,

Director, Antimicrobials Division, Office of Pesticide Programs.

[FR Doc. 2023–07727 Filed 4–12–23; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–10881–01–OA]

Local Government Advisory Committee (LGAC) and Small Communities Advisory Subcommittee (SCAS) Meeting

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notification of public meeting.

SUMMARY: Pursuant to the Federal Advisory Committee Act (FACA), EPA herby provides notice of a meeting for the Local Government Advisory Committee (LGAC) and its Small Communities Advisory Subcommittee (SCAS) on the date and time described below. This meeting will be open to the public. For information on public attendance and participation, please see the registration information under

DATES: The LGAC will have a hybrid meeting on May 22nd 2023, from 1 to 5 p.m. Eastern Daylight Time and May 23rd, 2023 from 8:30 a.m. to 12 p.m. Eastern Daylight Time. The SCAS will have a hybrid meeting prior to the LGAC on May 22nd, 2023, from 10:30 a.m. to 12 p.m. Eastern Daylight Time.

FOR FURTHER INFORMATION CONTACT: Paige Lieberman, Designated Federal Officer (DFO), at *LGAC@epa.gov* or 202–564–9957.

Information on Accessibility: For information on access or services for individuals requiring accessibility accommodations, please contact Paige Lieberman by email at *LGAC@epa.gov.* To request accommodation, please do so five (5) business days prior to the meeting, to give EPA as much time as possible to process your request.

SUPPLEMENTARY INFORMATION:

Content

The LGAC will discuss several priority issues at EPA, including providing draft recommendations on proposed national drinking water quality standards for PFAS, continuing discussions on climate mitigation, environmental justice and risk communications regarding PFAS. The SCAS will review these issues, as well as discuss recommendations on land use and transportation issues for small communities. Both the LGAC and SCAS will hear from EPA leadership regarding several new proposed charges. Details on the charges will be posted online (link below) one week prior to the meeting.

Registration

The meeting will be held virtually as well as in person. Members of the public who wish to participate should register by contacting the Designated Federal Officer (DFO) at *LGAC@epa.gov* by May 19, 2023. Online participation will be via Microsoft Teams. In person participation will be at EPA Headquarters, 1200 Constitution Ave. NW, Washington, DC.

Once available, the agenda and other supportive meeting materials will be available online at *https://www.epa.gov/ocir/local-government-advisory-committee-lgac* and will be emailed to all registered. In the event of cancellation for unforeseen circumstances, please contact the DFO or check the website above for reschedule information.

Dated: April 3, 2023.

Paige Lieberman,

Designated Federal Officer, U.S. Environmental Protection Agency.

[FR Doc. 2023–07758 Filed 4–12–23; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality; Notice of Meetings

AGENCY: Agency for Healthcare Research and Quality (AHRQ), Department of Health and Human Services (HHS).

ACTION: Notice of Five AHRQ Subcommittee Meetings.

SUMMARY: The subcommittees listed below are part of AHRQ's Health Services Research Initial Review Group (IRG) Committee. Grant applications are to be reviewed and discussed at these meetings. Each subcommittee meeting will be closed to the public.

DATES: See below for dates of meetings:

1. *Healthcare Research Training (HCRT)*
   Date: May 18–19, 2023
   July 14, 2023
2. *Healthcare Safety and Quality Improvement Research (HSQR)*
   Date: The date of the HSQR meeting is yet to be determined and will be published in an upcoming notice.
3. *Healthcare Information Technology Research (HITR)*
   Date: June 1–2, 2023
4. *Healthcare Effectiveness and Outcomes Research (HEOR)*
   Date: June 7–8, 2023
5. *Health System and Value Research (HSVR)*
   Date: June 15–16, 2023

ADDRESSES: Agency for Healthcare Research and Quality (Virtual Review for HCRT, HEOR & HSVR) 5600 Fishers Lane, Rockville, Maryland 20857, *and* Bethesda North Marriott Hotel & Conference Center (HITR in person review), 5701 Marinelli Road, Rockville, MD 20852–2785.

FOR FURTHER INFORMATION CONTACT: (to obtain a roster of members, agenda or minutes of the non-confidential portions of the meetings.)

Jenny Griffith, Committee Management Officer, Office of Extramural Research Education and Priority Populations, Agency for Healthcare Research and Quality (AHRQ), 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 427–1557.

SUPPLEMENTARY INFORMATION: In accordance with section 10 (a)(2) of the Federal Advisory Committee Act (5 U.S.C. App. 2), AHRQ announces meetings of the above-listed scientific peer review groups, which are subcommittees of AHRQ's Health Services Research Initial Review Group Committee. The subcommittee meetings will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2 section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Agenda items for these meetings are subject to change as priorities dictate.

Dated: April 7, 2023.

Marquita Cullom,

Associate Director.

[FR Doc. 2023–07747 Filed 4–12–23; 8:45 am]

BILLING CODE 4160–90–P