This Draft CII General Permit is established pursuant to CWA sections 301, 402(a)(1), 402(p)(2)(E), and 402(p)(6).

1. Technology-Based Requirements

This Draft CII General Permit would be available to authorize certain stormwater discharges, as it is authorized to do under CWA sections 301, 402(a)(1), 402(p)(2)(E) and 402(p)(6), by requiring the implementation of BMPs. The Draft CII Permit regulates Phosphorus, a non-conventional pollutant, as an indicator parameter for all regulated pollutants. As provided in section 402(a)(1) of the CWA, EPA established Technology-Based Effluent Limitations (TBELs) in this Draft Permit utilizing Best Professional Judgment (BPJ) to meet the ''best available technology economically achievable'' (BAT), ''best conventional pollutant control technology'' (BCT), and ''best practicable control technology currently available'' (BPT) standards described in section 304(b) of the CWA. TBELs in this Draft Permit are expressed as requirements for implementation of effective best management practices (BMPs). 40 CFR 122.44(k). Section 2.1.1 of the Draft CII GP requires all Permittees to develop and implement Stormwater Pollution Control Plans (SPCPs). The minimum BMPs specified in this CII GP represent common practices that can be implemented by most CII facilities. Dischargers have flexibility in designing their SPCP in accordance with Section 2.2 of this Draft CII GP.

2. Water-Quality Based Requirements

Based on its scientific and technical judgment, EPA has determined that the following reductions of Phosphorus, as an indicator pollutant, from CII sites are necessary to meet water quality standards: Charles River watershed, 65%; Mystic River watershed, 62%; Neponset River watershed, 60%. Permittees' development and implementation of SPCPs constitutes compliance with the Water-Quality Based Effluent Limitations (WQBELs) contained in this Draft CII GP, including the aforementioned reductions of Phosphorus.

D. Provisions on Which EPA Is Soliciting Comment

While EPA encourages the public to review and comment on all provisions in the Preliminary Determination and the Draft CII GP, EPA has included in the body of the Draft CII GP Fact Sheet several proposed provisions on which EPA specifically requests feedback. The following list summarizes these specific requests for comment, and where they are included in the fact sheet. EPA notes that these are only summaries of the requests for comment. The Agency recommends that the public see the specific wording of each comment request within the body of the fact sheet.

1. *Multifamily housing/tax codes (Fact Sheet § 1.5):* EPA is seeking comment on whether to include Multi-Family Residential Properties in the final designation and in the final CII GP.

2. *Compliance schedule (Fact Sheet § 5.1.1):* EPA is seeking comment on whether the proposed compliance schedule is appropriate.

3. *Multiple non-contiguous properties (Fact Sheet § 1.4):* EPA is seeking comment on how the permitting process should work for owners with multiple non-contiguous properties that are subject to the CII GP.

4. *Owner-operator (Fact Sheet § 1.4):* EPA is seeking comment on whether EPA should regulate the operator with control over a site instead of the owner, including sites where multiple operators may be tenants of a site (*e.g.,* a shopping plaza with one owner and multiple tenants).

5. *Contiguous properties (Fact Sheet § 1.4):* EPA is seeking comment on its regulation of contiguous sites, which reflect EPAs interest in consolidating, to the greatest extent possible, responsibility for permit compliance.

6. *Historic Properties (Fact Sheet § 9.3):* EPA is seeking comment on the Draft CII GP's potential impact on historic properties.

E. Procedures for Reaching a Final Designation and Final Permit Decision

After the comment period closes, EPA intends to issue a final permit and final RDA determination. EPA will consider all significant comments and make appropriate changes before issuing this permit. EPA's responses to public comments received will be included in the docket as part of the final permit issuance. Once the final permit becomes effective, eligible dischargers may seek authorization.

Authority: This action is being taken pursuant to Clean Water Act sections 301, 402(a)(1), 402(p)(2)(E), and 402(p)(6).

Dated: October 24, 2024.

David W. Cash,

Administrator, EPA Region 1, Boston, MA.

[FR Doc. 2024–25219 Filed 10–30–24; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL ELECTION COMMISSION

Sunshine Act Meetings

FEDERAL REGISTER CITATION OF PREVIOUS ANNOUNCEMENT: 89 FR 85212.

PREVIOUSLY ANNOUNCED TIME AND DATE OF THE MEETING: Wednesday, October 30, 2024 at 10:00 a.m., Hybrid Meeting: 1050 First Street NE, Washington, DC (12th Floor) and virtual.

CHANGES IN THE MEETING: The October 30, 2024 Open Meeting has been canceled.

CONTACT PERSON FOR MORE INFORMATION: Judith Ingram, Press Officer, Telephone: (202) 694–1220.

(Authority: Government in the Sunshine Act, 5 U.S.C. 552b)

Laura E. Sinram,

Secretary and Clerk of the Commission.

[FR Doc. 2024–25498 Filed 10–29–24; 4:15 pm]

BILLING CODE 6715–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Meeting on the Artificial Intelligence in Healthcare Safety Program

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

ACTION: Notice of public meeting.

SUMMARY: HHS is directed by Executive order (E.O.) to establish an Artificial Intelligence (AI) in Healthcare Safety Program in partnership with federally listed Patient Safety Organizations (PSOs). The purpose of this notice is to announce a meeting to discuss implementation of the Executive order to establish the AI in Healthcare Safety Program. This meeting is designed as an interactive forum where participants can provide input on the future of the program.

DATES: The meeting will be held from 12:30 to 4 p.m. eastern on Friday, November 15, 2024.

ADDRESSES: The meeting will be held virtually.

FOR FURTHER INFORMATION CONTACT: Erofile Gripiotis, Program Analyst, Center for Quality Improvement and Patient Safety, AHRQ, 5600 Fishers Lane, Rockville, MD 20857; Telephone (toll free): (866) 403–3697; Telephone (local): (301) 427–1111; TTY (toll free): (866) 438–7231; TTY (local): (301) 427–1130; Email: *pso@ahrq.hhs.gov*.

SUPPLEMENTARY INFORMATION:

Background

The Patient Safety and Quality Improvement Act of 2005, 42 U.S.C. 299b–21 to 299b–26 (Patient Safety Act), and the related Patient Safety and Quality Improvement Final Rule, 42 CFR part 3 (Patient Safety Rule), published in the **Federal Register** on November 21, 2008, 73 FR 70731–70814, provide for the Federal listing of Patient Safety Organizations (PSOs), which collect, aggregate, and analyze confidential information (patient safety work product) regarding the quality and safety of healthcare delivery.

The Patient Safety Act also authorizes the development of data standards, known as the Common Formats, to facilitate the aggregation and analysis of non-identifiable patient safety work product collected by PSOs and reported to the Network of Patient Safety Databases (NPSD). (42 U.S.C. 299b–23(b)). The Patient Safety Act and Patient Safety Rule can be accessed at: *http://www.pso.ahrq.gov/legislation/*.

Section 8(iv) of E.O. 14110 requires the Secretary of HHS, in consultation with the Secretaries of Defense and Veterans Affairs, to establish an AI safety program that, in partnership with PSOs, will:

- establish a common framework for approaches to identifying and capturing clinical errors resulting from AI deployed in healthcare settings as well as specifications for a central tracking repository for associated incidents that cause harm, including through bias or discrimination, to patients, caregivers, or other parties;
- analyze captured data and generated evidence to develop, wherever appropriate, recommendations, best practices, or other informal guidelines aimed at avoiding these harms; and
- disseminate those recommendations, best practices, or other informal guidelines to appropriate stakeholders, including healthcare providers.

Agenda, Registration, and Other Information About the Meeting

AHRQ will be hosting this fully virtual meeting to discuss implementation of the AI in Healthcare Safety Program with members of the public, including PSOs and other interested parties. Agenda topics will include recent AI-related analyses from the NPSD, available program resources, and speakers from the Assistant Secretary for Technology Policy/Office of the National Coordinator for Health Information Technology and the Coalition for Health AI. Active participation and discussion by meeting participants is encouraged, including through breakout sessions.

AHRQ requests that interested persons send an email to *SDMeetings@infinityconferences.com* for registration information. Before the meeting, an agenda and logistical information will be provided to registrants.

Dated: October 24, 2024.

Marquita Cullom,

Associate Director.

[FR Doc. 2024–25140 Filed 10–30–24; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC–2024–0065; Docket Number NIOSH–352–A]

Draft Hazard Review: Wildland Fire Smoke Exposure Among Farmworkers and Other Outdoor Workers

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Extension of comment period and announcement of informational webinar.

SUMMARY: On September 13, 2024, NIOSH published a notice in the **Federal Register** announcing public comment and technical review on the draft Hazard Review: Wildland Fire Smoke Exposure Among Farmworkers and Other Outdoor Workers. Written comments were to be received by November 12, 2024. NIOSH is extending the public comment period to January 10, 2025. NIOSH will also convene an informational webinar to present an overview about the draft Hazard Review document, describe its content and purpose, and provide information about the public comment period. The webinar is scheduled to occur on Tuesday, December 3, 2024, at 1:00 p.m. Eastern Time (US and Canada). Attendees are requested to register in advance for this webinar.

DATES: Registration for the webinar must occur on or before the date of the webinar, December 3, 2024. The comment period for the draft Hazard Review: Wildland Fire Smoke Exposure Among Farmworkers and Other Outdoor Workers, published September 13, 2024 at 89 FR 74960, is extended. Electronic or written comments must be received by January 10, 2025, at 11:59 p.m.

ADDRESSES: Register in advance for the webinar at the following link. Attendance for the webinar is limited to 3,000 participants: *https://cdc.zoomgov.com/webinar/register/WN_lnOhz1wMTNyr_06z1hYgRw.*

You may submit comments on the draft Hazard Review: Wildland Fire Smoke Exposure Among Farmworkers and Other Outdoor Workers, identified by CDC–2024–0065 and Docket Number NIOSH–352–A, by either of the following two methods:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C–34, Cincinnati, Ohio 45226–1998.

Instructions: All information received in response to this notice must include the agency name and docket number (CDC–2024–0065; NIOSH–352–A). All relevant comments, including any personal information provided, will be posted without change to *https://www.regulations.gov.* Do not submit comments by email. CDC does not accept comments by email. For access to the docket to read the draft Hazard Review document or comments received, go to *https://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: R. Todd Niemeier, Ph.D., National Institute for Occupational Safety and Health, MS–C15, 1090 Tusculum Avenue, Cincinnati, OH 45226. Telephone: (513) 533–8166.

SUPPLEMENTARY INFORMATION: NIOSH is requesting public comment and technical review of the draft Hazard Review: Wildland Fire Smoke Exposure Among Farmworkers and Other Outdoor Workers, which is accessible in the docket (CDC–2024–0065; NIOSH–352–A). NIOSH is extending the public comment period to January 10, 2025. The comment period is being extended to provide an informational webinar and allow additional time for comment. Specific review questions to be considered are included in the initial **Federal Register** notice published on September 13, 2024 at 89 FR 74960.

The final document will be used as the scientific evidence base to inform the development of supplementary educational materials for workers, employers, and other relevant audiences to support the implementation of the recommendations. Therefore, comments that focus on the understandability, accessibility, and feasibility of the recommendations are requested.

The draft Hazard Review was developed to provide the scientific rationale for characterizing hazards of exposure to wildland fire smoke for