III. What is the DfE program?

The DfE program is a companion program to Safer Choice and certifies antimicrobial products. The DfE logo may be used on certified products and helps consumers and commercial buyers identify products that meet the health and safety standards of the pesticide registration process required by the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) as well as the Safer Choice program's stringent criteria for efficacy and effects on human health and the environment. For more information on the DfE program, please see: *https://www.epa.gov/pesticide-labels/learn-about-design-environment-dfe-certification.*

IV. What is the purpose of the award?

The purpose of the Partner of the Year Awards is to recognize the leadership contributions of Safer Choice program partners and stakeholders who, over the past year, have shown achievement in the design, manufacture, promotion, selection, and use of products with safer chemicals, that further outstanding or innovative source reduction. EPA especially encourages submission of award applications that show how the applicant's work involving products with safer chemical ingredients promotes environmental justice, bolsters resilience to the impacts of climate change, results in cleaner air or water, improves drinking water quality, or advances innovation in packaging. Similar achievement in the design, manufacture, promotion, selection, and use of Design for the Environment (DfE)-certified products will also make an organization eligible for the Partner of the Year Awards.

V. How can I participate?

All Safer Choice stakeholders and program participants in good standing are eligible for recognition. Interested parties who would like to be considered for this award should submit to EPA an application detailing their accomplishments and contributions during calendar year 2023. The application form is available on the Safer Choice website. Candidates interested in learning more about the Partner of the Year Awards should refer to the following link: *https://www.epa.gov/saferchoice/safer-choice-partner-year-awards.* EPA will recognize award winners at a Safer Choice Partner of the Year Awards ceremony later in 2024.

Authority: 42 U.S.C. 13103(b)(13) and 15 U.S.C. 2609.

Dated: January 26, 2024.

Michal Freedhoff,

Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

[FR Doc. 2024–01909 Filed 1–30–24; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OAR–2023–0137; FRL–11708–01–OMS]

Agency Information Collection Activities; Submission to the Office of Management and Budget for Review and Approval; Comment Request; New Source Performance Standards (NSPS) for Sewage Sludge Incineration Units (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted an information collection request (ICR), NSPS for Sewage Sludge Incineration Units (EPA ICR Number 2369.06, OMB Control Number 2060–0658), to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act. This is a proposed extension of the ICR, which is currently approved through January 31, 2024. Public comments were previously requested, via the **Federal Register** on May 18, 2023, during a 60-day comment period. This notice allows for an additional 30 days for public comments.

DATES: Comments may be submitted on or before March 1, 2024.

ADDRESSES: Submit your comments, referencing Docket ID Number EPA–HQ–OAR–2023–0137, to EPA online using *www.regulations.gov/* (our preferred method), or by email to *a-and-r-docket@epa.gov,* or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW, Washington, DC 20460. The EPA's policy is that all comments received will be included in the public docket without change, including any personal information provided, unless the comment includes profanity, threats, information claimed to be Confidential Business Information (CBI), or other information whose disclosure is restricted by statute.

Submit written comments and recommendations to OMB for the proposed information collection within 30 days of publication of this notice to: *www.reginfo.gov/public/do/PRAMain.* Find this specific information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Muntasir Ali, Sector Policies and Program Division (D243–05), Office of Air Quality Planning and Standards, U.S. Environmental Protection Agency, Research Triangle Park, North Carolina, 27711; telephone number: (919) 541–0833; email address: *ali.muntasir@epa.gov.*

SUPPLEMENTARY INFORMATION: This is a supporting extension of this ICR, which is currently approved through January 31, 2024. An agency may neither conduct nor sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

Public comments were previously requested, via the **Federal Register**, on July 22, 2022, during a 60-day comment period (87 FR 43843) and May 18, 2023 (88 FR 31748). This notice allows for an additional 30 days for public comments. Supporting documents, which explain in detail the information that the EPA will be collecting, are available in the public docket for this ICR. The docket can be viewed online at *www.regulations.gov,* or in person at the EPA Docket Center, WJC West Building, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The telephone number for the Docket Center is 202–566–1744. For additional information about EPA's public docket, visit *http://www.epa.gov/dockets.*

Abstract: The New Source Performance Standards for Sewage Sludge Incineration Units (40 CFR part 60, subpart LLLL) were proposed on October 14, 2010; and promulgated on March 21, 2011. These regulations apply to new facilities with one or more sewage sludge incineration (SSI) units. New facilities are those that either commenced construction after October 14, 2010, or commenced modification after September 21, 2011. Physical or operational changes made to the SSI unit to comply with the SSI Emission Guidelines at 40 CFR part 60, subpart MMMM do not qualify as a modification under this NSPS. This information is being collected to assure compliance with 40 CFR part 60, subpart LLLL.

Form Numbers: None.

Respondents/affected entities: Sewage Sludge Incinerators.

Respondent's obligation to respond: Mandatory (40 CFR part 60, subpart LLLL).

Estimated number of respondents: 11 (total).

Frequency of response: Semiannual, annual.

Total estimated burden: 1,800 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated cost: $1,950,000 (per year), which includes $1,850,000 in annualized capital/startup and/or operation & maintenance costs.

Changes in the Estimates: The increase in burden from the most-recently approved ICR is due to an increase in the number of new or modified sources. There is also an increase in costs due to the use of updated labor rates. This ICR uses labor rates from the most-recent Bureau of Labor Statistics report (September 2022) to calculate respondent burden costs. This ICR also adjusts the capital/startup and operation and maintenance costs from 2008 to 2022 values using the CEPCI CE Index.

Courtney Kerwin,

Director, Regulatory Support Division.

[FR Doc. 2024–01807 Filed 1–30–24; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10788]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by April 1, 2024.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number ____: Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10788 Prescription Drug and Health Care Spending

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. Type of Information Collection Request: Revision of a currently approved collection; *Title of Information Collection:* Prescription Drug and Health Care Spending; *Use:* On December 27, 2020, the Consolidated Appropriations Act, 2021 (CAA) was signed into law. Section 204 of title II of division BB of the CAA added parallel provisions at section 9825 of the Internal Revenue Code (the Code), section 725 of the Employee Retirement Income Security Act (ERISA), and section 2799A–10 of the Public Health Service Act (PHS Act) that require group health plans and health insurance issuers offering group or individual health insurance coverage to annually report to the Department of the Treasury, the Department of Labor (DOL), and the Department of Health and Human Services (HHS) (collectively, ''the Departments'') certain information about prescription drug and health care spending, premiums, and enrollment under the plan or coverage. This information will support the development of public reports that will be published by the Departments on prescription drug reimbursements for plans and coverage, prescription drug pricing trends, and the role of prescription drug costs in contributing to premium increases or decreases under the plans or coverage. The 2021 interim final rules, ''Prescription Drug and Health Care Spending'' (2021 interim final rules), issued by the Departments and the Office of Personnel Management (OPM) implement the provisions of section 9825 of the Code, section 725 of ERISA, and section 2799A–10 of the PHS Act, as enacted by section 204 of title II of division BB of the CAA. OPM joined the Departments in issuing the 2021 interim final rules, requiring Federal Employees Health Benefits (FEHB) carriers to report information about prescription drug and health care spending, premiums, and plan enrollment in the same manner as a group health plan or health insurance issuer offering group or individual health insurance coverage.

The 2023 Prescription Drug Data Collection (RxDC) Reporting Instructions reflect changes for the 2023 reference year and beyond. As a result of removing one-time first- and second-