Estimated number of respondents: 56 (total).

Frequency of response: Initially, occasionally, and semiannually.

Total estimated burden: 28,700 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated cost: $3,010,000 (per year), includes $0 annualized capital or operation & maintenance costs.

Changes in the Estimates: There is an increase of 106 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. This adjustment in the burden occurred because this ICR assumes all existing respondents will have to familiarize with the regulatory requirements each year.

Courtney Kerwin, Director, Regulatory Support Division.

ADDRESS: Submit your comments, referencing Docket ID Number EPA–HQ–OECA–2013–0324, to: (1) EPA online using www.regulations.gov (our preferred method), or by email to docket.oeca@epa.gov, or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW, Washington, DC 20460; and (2) OMB via email to oira_submission@omb.eop.gov. Address comments to OMB Desk Officer for EPA.

EPA’s policy is that all comments received will be included in the public dockets without changes including any personal information provided, unless the comment includes profanity, threats, or information claimed to be Confidential Business Information (CBI), or other information whose disclosure is restricted by statute.

FOR FURTHER INFORMATION CONTACT: Patrick Yellin, Monitoring, Assistance, and Media Programs Division, Office of Compliance, Mail Code 2227A, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: (202) 564–2970; fax number: (202) 564–0050; email address: yellin.patrick@epa.gov.

SUPPLEMENTARY INFORMATION:

Supporting documents, which explain in detail the information that the EPA will be collecting, are available in the public dockets for this ICR. The docket can be viewed online at www.regulations.gov or in person at the EPA Docket Center, WJC West, 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 dockets, visit: http://www.epa.gov/dockets.

Abstract: The National Emission Standards for Hazardous Air Pollutants (NESHAP) for Marine Tank Vessel Loading Operations (40 CFR part 63, subpart Y) establishes Maximum Achievable Control Technology (MACT) standards for existing facilities and new facilities that load marine tank vessels with petroleum or gasoline. These facilities have aggregate actual hazardous air pollutants (HAP) emissions of 10 tons or more of each individual HAP, or 25 tons or more of all HAP combined. This NESHAP regulation also established reasonably-available control technology (RACT) standards to such facilities with an annual throughput of 10 million or more barrels of gasoline or 200 million or more barrels of crude oil. The NESHAP regulation was amended in 2011 to include emission standards for two marine tank vessel loading operation (MTVLO) subcategories not included in the original rule. These subcategories are facilities with MTVLO that emit less than 10 tons per year of each individual HAP and less than 25 tons per year of all HAP combined, and that are located at major sources of HAP loading more than 1 million barrels per year of gasoline, and facilities located more than 0.5 miles from shore. The 2011 amendment also added a provision to require electronic submittal of performance test results. This ICR has been updated to reflect the additional industry burden associated with the amended standards. The 2015 amendments did not add information collection requirements beyond those currently required under the applicable regulations.

In general, all NESHAP standards require initial notifications, performance tests, and periodic reports by the owners/operators of the affected facilities. They are also required to maintain records of the occurrence and duration of any startup, shutdown, or malfunction in the operation of an affected facility or any period during which the monitoring system is inoperative. These notifications, reports, and records are essential in determining compliance, and are required of all affected facilities subject to NESHAP. Any owner/operator subject to the provisions of this part shall maintain a file of these measurements, and retain the file for at least five years following the date of such measurements, maintenance reports, and records. All reports are sent to the delegated state or local authority. If there is no such delegated authority, the reports are sent directly to the EPA regional office.

Respondent’s obligation to respond: Mandatory (40 CFR part 63, subpart Y).

Estimated number of respondents: 804 (total).

Frequency of response: Initially and annually.

Total estimated burden: 10,700 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated cost: $1,130,000 (per year), which includes no annualized capital/startup or operation & maintenance costs.

Changes in the Estimates: There is a small increase in the total estimated burden as currently identified in the OMB Inventory of Approved Burdens. This adjustment is due to a change in a previous assumption. In accordance with the Terms of Clearance for OMB’s previous approval, this ICR renewal
DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers CMS–10143 and CMS–10516]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services.

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 (the 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 June 19, 2018.

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, you may make your request using one of following:


2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: William 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–10143 State Data for the Medicare Modernization Act (MMA)

CMS–10516 Program Integrity: Exchange, Premium Stabilization Programs, and Market Standards; Amendments to the HHS Notice of Benefit and Payment Parameters for 2014; Final Rule II

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: State Data for the Medicare Modernization Act (MMA); Use: The monthly data file is provided to CMS by states on dual eligible beneficiaries. The phase-down process requires a monthly count of all full benefit dual eligible beneficiaries with an active Part D plan enrollment in the month. CMS will make this selection of records using dual eligibility status codes contained in the person-month record to identify all full-benefit dual eligible beneficiaries (codes 02, 04 and 08). In the case where in a given month, multiple records were submitted for the same beneficiary in multiple file submittals, the last record submitted for that beneficiary shall be used to determine the final effect on the phase-down count. Form Number: CMS–10143 (OMB Control Number: 0938–0958); Frequency: Monthly; Affected Public: State, Local, or Tribal Governments; Number of Respondents: 51; Total Annual Responses: 612; Total Annual Hours: 4,896. (For policy questions regarding this collection contact Linda King at 410–786–1312.)

2. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Program Integrity: Exchange, Premium Stabilization Programs, and Market Standards; Amendments to the HHS Notice of Benefit and Payment Parameters for 2014; Final Rule II; Use: The original approved ICR affiliated with this final rule (OMB #: 0938–1277) was titled Program Integrity: Exchange, Premium Stabilization Programs, and Market Standards; Amendments to the HHS Notice of Benefit and Payment Parameters for 2014; Final Rule II and was approved on 8/26/2015. This Information Collection Request (ICR) serves as the formal request for renewal of the clearance. This ICR includes some of the ICRs from the previously approved final rule. The program integrity data collections and third-party disclosure requirements will assist HHS in determining Exchange compliance with Federal standards. The data collection and third-party disclosure requirements will also assist HHS in monitoring QHP issuers in FFEs for compliance with Federal QHP issuer standards. The data collected by health insurance issuers and Exchanges will