C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the Regulatory Secretariat Division by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.* Please cite OMB Control No. 3090–0027, Contract Administration, Quality Assurance (GSA Forms 1678 and 308), in all correspondence.

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2024–20387 Filed 9–9–24; 8:45 am]

BILLING CODE 6820–61–P

OFFICE OF GOVERNMENT ETHICS

Agency Information Collection Activities; Submission for OMB Review; Information Collection Renewal; Comment Request; Fast Track Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

AGENCY: Office of Government Ethics (OGE).

ACTION: Notice of request for comments.

SUMMARY: After this second round notice, the U.S. Office of Government Ethics (OGE) intends to submit a request for a renewed Generic Clearance for the collection of qualitative feedback on agency service delivery for review and approval of a three-year extension under the Paperwork Reduction Act.

Comments: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Jennifer Matis at the U.S. Office of Government Ethics; telephone: 202–482–9216; TTY: 800–877–8339; Email: *usoge@oge.gov.*

SUPPLEMENTARY INFORMATION:

Title: Fast Track Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

Abstract: The proposed Generic Clearance provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the agency's commitment to improving service delivery. Qualitative feedback means information that provides useful insights on perceptions and opinions but is not a statistical survey that yields quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences, and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training, or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative, and actionable communications between the agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

OGE expects to use various methods (*e.g.,* focus groups, customer satisfaction surveys, comment cards) to solicit feedback. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public and other agency stakeholders. If this information is not collected, vital feedback from customers and stakeholders on the agency's services will be unavailable.

The agency will only submit a collection for approval under this Generic Clearance if it meets the following conditions:

- The collections are voluntary;
- The collections are low-burden for respondents (based on considerations of total burden hours, total number of respondents, or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government;
- The collections are noncontroversial;
- The collections are focused on the awareness, understanding, attitudes, preferences, or experiences of the public or other stakeholders in order to improve existing or future services, products, or communication materials;
- Personally identifiable information (PII) is collected only to the extent necessary;
- Information gathered will be used only internally for general service improvement and program management purposes and is not intended for release to the public;
- Information gathered will not be used for the purpose of substantially informing influential policy decisions; and
- Information gathered will yield qualitative information, and the collections will not be designed or expected to yield statistically reliable results or used as though the results are generalizable to the population of study.

Feedback collected under this Generic Clearance provides useful information, but it does not yield data that can be generalized to the overall population. This type of Generic Clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: the target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential nonresponse bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior to fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

As a general matter, information collections submitted under this Generic Clearance will not result in any new system of records containing privacy information and will not ask questions of a sensitive nature, such as sexual behavior and attitudes, religious beliefs, and other matters that are commonly considered private.

A **Federal Register** Notice with a 60-day comment period soliciting comments on this information collection was published on June 17, 2024 (89 FR 51344). OGE received no comments.

OMB Number: 3209–0010.

Type of Request: Extension.

Affected Public: Individuals; Business or Other For-Profit Institutions; Not-For-Profit Institutions; State, Local, or Tribal Government.

Estimated Annual Number of Respondents: 91,555.

Average Expected Annual Number of Activities: 6.

Average Number of Respondents per Activity: 15259.

Responses per Respondent: 1.

Annual Responses: 91,555.

Average Minutes per Response: 56 minutes.

Annual Burden Hours: 4,030 hours.

Frequency: On occasion.

Request for Comments: Public comment is invited specifically on the need for and practical utility of this Generic Clearance, the accuracy of OGE's burden estimate, the enhancement of quality, utility and clarity of the information collected, and the minimization of burden (including the use of information technology). The comments will become a matter of public record.

Approved: September 4, 2024.

Shelley K. Finlayson,

Acting Director, U.S. Office of Government Ethics.

[FR Doc. 2024–20352 Filed 9–9–24; 8:45 am]

BILLING CODE 6345–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10141, CMS–10913, CMS–R–290 and CMS–10443]

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 November 12, 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–10141 Medicare Prescription Drug Benefit Program

CMS–10913 Medicare Part C Utilization Management Annual Data Submission and Audit Protocol Data Request

CMS–R–290 Medicare Program: Procedures for Making National Coverage Decisions

CMS–10443 Transcatheter Valve Therapy (TVT) Registry

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 Collections

1. *Type of Information Collection Request:* Revision of a currently approved information collection; *Title of Information Collection:* Medicare Prescription Drug Benefit Program; *Use:* CMS will use this information from plan sponsors and States to approve contract applications, monitor compliance with contract requirements, make proper payment to plans, and ensure that correct information is disclosed to potential and current enrollees. *Form Number:* CMS–10141 (OMB control number: 0938–0964); *Frequency:* Annually; *Affected Public:* Private Sector, State, Local, or Tribal Governments; *Number of Respondents:* 4,633,032; *Total Annual Responses:* 87,014,803; *Total Annual Hours:* 25,409,037. (For policy questions regarding this collection contact Chad Buskirk at 410–786–1630 or *chad.buskirk@cms.hhs.gov*).

2. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Medicare Part C Utilization Management Annual Data Submission and Audit Protocol Data Request; *Use:* Section 1857(d) of the Act, added by the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 and implementing regulations at 42 CFR 422.503 and 422.504 state that CMS must oversee an MA organization's continued compliance with the requirements for a MA organization. Additionally, per § 422.516(a), MA organizations are required to compile and report to CMS information related to the utilization of services, and other matters as CMS may require.

The information gathered during this annual data collection and audit will be used by the Medicare Parts C and D Oversight and Enforcement Group (MOEG) within the Center for Medicare (CM) to assess Sponsoring organizations' compliance with Medicare UM requirements. CMS will utilize the data submitted during the annual data submission to assess the number of items and services that have associated internal coverage criteria, and to develop a landscape of items and services across the nation to assess