Only intervenors have the right to seek rehearing of the Commission's decision and be heard by the courts if they choose to appeal the Commission's final ruling. An intervenor formally participates in the proceeding by filing a request to intervene pursuant to Rule 214 of the Commission's Rules of Practice and Procedures (18 CFR 385.214). Motions to intervene are more fully described at *https://www.ferc.gov/resources/guides/how-to.asp.* Please note that the Commission will not accept requests for intervenor status at this time. You must wait until the Commission receives a formal application for the project, after which the Commission will issue a public notice that establishes an intervention deadline.

Additional Information

Additional information about the project is available from the Commission's Office of External Affairs, at (866) 208–FERC, or on the FERC website (*www.ferc.gov*) using the eLibrary link. Click on the eLibrary link, click on ''General Search'' and enter the docket number in the Docket Number field. Be sure you have selected an appropriate date range. For assistance, please contact FERC Online Support at *FercOnlineSupport@ferc.gov* or toll free at (866) 208–3676, or for TTY, contact (202) 502–8659. The eLibrary link also provides access to the texts of formal documents issued by the Commission, such as orders, notices, and rulemakings.

Public meetings or site visits will be posted on the Commission's calendar located at *https://www.ferc.gov/news-events/events* along with other related information.

Dated: May 7, 2025.

Carlos D. Clay,

Deputy Secretary.

[FR Doc. 2025–08365 Filed 5–12–25; 8:45 am]

BILLING CODE 6717–01–P

EXECUTIVE OFFICE OF THE PRESIDENT

Office of National Drug Control Policy

Appointment of Members of Senior Executive Service Performance Review Board

AGENCY: Office of National Drug Control Policy (ONDCP).

ACTION: Notice of appointments.

SUMMARY: The following persons have been appointed to the ONDCP Senior Executive Service Performance Review Board: Ms. Michele Marx (as Chair), Ms. Debbie Seguin, Ms. Shannon Kelly, and Mr. Kemp Chester.

FOR FURTHER INFORMATION CONTACT: Please direct any questions to Anthony Jones, Acting General Counsel, (202) 881–8862, Office of National Drug Control Policy, Executive Office of the President, Washington, DC 20503.

Authority: 5 U.S.C. 4314(c)(1)

Dated: May 8, 2025.

Anthony Jones,

Acting General Counsel.

[FR Doc. 2025–08404 Filed 5–12–25; 8:45 am]

BILLING CODE 3280–F5–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–R–138 and CMS–10882]

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 July 14, 2025.

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–R–138 Medicare Geographic Classification Review Board Procedures and Criteria

CMS–10882 Part C and Part D Medicare Prescription Payment Plan Model Documents

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:* Reinstatement with change of a previously approved collection; *Title of Information Collection:* Medicare Geographic Classification Review Board Procedures and Criteria; *Use:* During the first few years of IPPS, hospitals were paid strictly based on their physical geographic location concerning the wage index (Metropolitan Statistical Areas (MSAs)) and the standardized amount (rural, other urban, or large urban). However, a growing number of hospitals became concerned that their payment rates were not providing accurate compensation. The hospitals argued that they were not competing with the hospitals in their own geographic area, but instead that they were competing with hospitals in neighboring geographic areas.

At that point, Congress enacted Section 1886(d)(10) of the Act which enabled hospitals to apply to be considered part of neighboring geographic areas for payment purposes based on certain criteria. The application and decision process are administered by the MGCRB which is not a part of CMS so that CMS could not be accused of any untoward action. However, CMS needs to remain apprised of any potential payment changes. Hospitals are required to provide CMS with a copy of any applications that they made to the MGCRB. CMS also developed the guidelines for the MGCRB that were the interim final issue of the **Federal Register** and must ensure that the MGCRB properly applied the guidelines. This check and balance process also contributes to limiting the number of hospitals that ultimately need to appeal their MGCRB decisions to the CMS Administrator. *Form Number:* CMS–R–138 (OMB control number: 0938–0573); *Frequency:* Occasionally; *Affected Public:* Businesses or other for-profits and Not-for-profit institutions; *Number of Respondents:* 850; *Total Annual Responses:* 850; *Total Annual Hours:* 850. (For policy questions regarding this collection contact Noel Manlove at 410–786–5161.)

2. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Part C and Part D Medicare Prescription Payment Plan Model Documents; *Use:* Sections 1860D–2(b)(2)(E)(v)(II)–(IV) of the Act state the requirements for Part D plan sponsors in implementing the program, which include the processes for outreach to enrollees identified as likely to benefit, election, and termination. Subsection II states that any Part D enrollee may elect into the program prior to or during the plan year. Subsection III details that Part D plan sponsors and MA organizations must have a mechanism in place to inform enrollees that they are likely to benefit from electing into the program at the point of sale (POS). Subsection IV(aa) states that plans must terminate a beneficiary's participation in the program when the beneficiary fails to pay the amounts owed under this program.

CMS has developed the seven model notices to provide standardized and consistent language for potential and active program participants, regardless of which Part D plan they may be enrolled in. The seven model notices and their content serve as an example of how to convey information on the Medicare Prescription Payment Plan to Part D enrollees and program participants, as applicable. Though Part D plan sponsors are not required to use the model materials and content verbatim, use of the model materials will satisfy the communications requirements included in § 423.137. If a Part D plan sponsor chooses not to use a model material, they must meet the content requirements in § 423.137 for the alternate notices they develop. CMS notes that the ''Medicare Prescription Payment Plan Likely to Benefit Notice,'' is a standardized material that Part D plan sponsors are required to use in the form and manner provided by CMS. *Form Number:* CMS–10882 (OMB control number: 0938–1475); *Frequency:* Yearly; *Affected Public:* Individuals and Households, Private Sector, Federal Government, Businesses or other for-profits and Not-for-profit institutions; *Number of Respondents:* 234,227; *Total Annual Responses:* 39,514,987; *Total Annual Hours:* 135,080. (For policy questions regarding this collection contact Deven Gosalia at (410) 786–8264 or *deven.gosalia@cms.hhs.gov*.)

William N. Parham, III,

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2025–08348 Filed 5–12–25; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10844]

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 July 14, 2025.

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 *https://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.