

GENERAL SERVICES ADMINISTRATION

[Notice—MA—2025—02; Docket No. 2024—0002, Sequence No. 51]

Calendar Year (CY) 2025 Privately Owned Vehicle (POV) Mileage Reimbursement Rates; CY 2025 Standard Mileage Rate for Moving Purposes

AGENCY: Office of Government-wide Policy (OGP), General Services Administration (GSA).

ACTION: Notice.

SUMMARY: GSA is updating the mileage reimbursement rate for privately owned automobiles (POA), airplanes, and motorcycles as required by statute. This information will be available in FTR Bulletin 25–04, which can be found on GSA’s website at <https://gsa.gov/ftbulletins>.

DATES: *Applicability date:* This notice applies to travel and relocation performed on or after January 1, 2025, through December 31, 2025.

FOR FURTHER INFORMATION CONTACT: For clarification of content, please contact Mrs. Autumn King, Policy Analyst, Office of Government-wide Policy, Office of Asset and Transportation Management, at 803–944–6487, or by email at travelpolicy@gsa.gov. Please cite Notice of FTR Bulletin 25–04.

SUPPLEMENTARY INFORMATION: GSA is required by statute to set the mileage reimbursement rate for privately owned automobiles (POA) as the single standard mileage rate established by the Internal Revenue Service (IRS). The IRS mileage rate for medical or moving purposes is used to determine the POA rate when a Government-furnished automobile is available and authorized and also represents the privately owned vehicle (POV) standard mileage reimbursement rate for official relocation.

Finally, GSA conducts independent reviews of the cost of travel and the operation of privately owned airplanes and motorcycles on an annual basis to determine their corresponding mileage reimbursement rates. These reviews evaluate various factors, such as the cost of fuel, depreciation of the original vehicle cost, maintenance and insurance, state and Federal taxes, and consumer price index data. FTR Bulletin 25–04 establishes and announces the new CY 2025 POV mileage reimbursement rates for official temporary duty and relocation travel.

This notice is the only notification to agencies of revisions to the POV mileage rates for official travel and relocation, in

addition to the changes posted on GSA’s website at <https://gsa.gov/mileage>.

Mehul Parekh,

Acting Associate Administrator, Office of Government-wide Policy.

[FR Doc. 2024–31556 Filed 1–2–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–209]

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 March 4, 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–209 Laboratory Personnel Report (CLIA) and Supporting Regulations

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 collection; *Title of Information Collection:* Laboratory Personnel Report (CLIA) and Supporting Regulations; *Use:* The information collected on this survey form is used in the administrative pursuit of the Congressionally-mandated program

with regard to regulation of laboratories participating in CLIA. The surveyor will provide the laboratory with the CMS–209 form. While the surveyor performs other aspects of the survey, the laboratory will complete the CMS–209 by recording the personnel data needed to support their compliance with the personnel requirements of CLIA. For this submission, we are making minor revisions to the collection instrument. We revised the instructions for clarity and removed the references to specific regulations. *Form Number:* CMS–209 (OMB control number 0938–0151); *Frequency:* Biennially; *Affected Public:* Private Sector—State, Local, or Tribal Governments; and Federal Government; *Number of Respondents:* 16,404; *Total Annual Responses:* 8,202; *Total Annual Hours:* 4,101. (For policy questions regarding this collection contact Penny Keller at 410–786–2035.)

William N. Parham, III,

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

[FR Doc. 2024–31553 Filed 1–2–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10565 and CMS–1763]

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 March 4, 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–10565 Medicare Advantage Model of Care Submission Requirements
CMS–1763 Request for Termination of Medicare Premium Part A, Part B, or Part B Immunosuppressive Drug Coverage (Part B–ID) and Supporting Statute and Regulations

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 collection; *Title of Information Collection:* Medicare Advantage Model of Care Submission Requirements; *Use:* Section 1859(f)(7) of the Act and 42 CFR 422.101(f)(3) requires that all SNP MOCs be approved by NCQA. This approval is based on NCQA's evaluation of SNPs' MOC narratives using MOC scoring guidelines. Section 50311 of the BBA of 2018 modified the MOC requirements for C–SNPs in section 1859 (f)(5)(B)(i–v) of the Act, requiring them to submit on an annual basis. The BBA mandated additional changes for C–SNPs related to care management, HRAs, individualized care plans, a minimum benchmark for scoring, etc., for which CMS has applied these requirements to all SNP types.

SNPs will submit initial and renewal MOCs as well as summaries of any substantive off-cycle MOC changes to CMS through HPMS. This is the platform that CMS uses to coordinate communication and the collection of information from MAOs.

NCQA and CMS will use information collected in the SNP Application HPMS module to review and approve MOC narratives in order for an MAO to offer a new SNP in the upcoming calendar year(s). This information is used by CMS as part of the MA SNP application process. NCQA and CMS will use information collected in the Renewal Submission section of the HPMS MOC module to review and approve the MOC narrative for the SNP to receive a new approval period and operate in the upcoming calendar year(s). *Form Number:* CMS–10565 (OMB control number 0938–1296); *Frequency:* Occasionally; *Affected Public:* Private Sector, Business or other for-profits; *Number of Respondents:* 2,088; *Total Annual Responses:* 2,088; *Total Annual Hours:* 8,638. (For policy questions regarding this collection contact Daniel