

*Estimated Time per Response:* 27.3–95 hours.

*Frequency of Response:*  
Recordkeeping requirement.

*Obligation to Respond:* Required to obtain or retain benefits. The statutory authority for this collection is contained in 47 U.S.C. 151–155, 301–609.

*Total Annual Burden:* 557,120 hours.

*Annual Cost Burden:* No cost.

*Needs and Uses:* The Commission will submit this extension (no change in the recordkeeping requirement) to the OMB after this 60 day comment period to obtain the full three-year clearance from them. The information collection requirements are as follows:

*Section 80.409(c), Public Coast Station Logs:* This requirement is necessary to document the operation and public correspondence of public coast radio telegraph, public coast radiotelephone stations, and Alaska public-fixed stations, including the logging of distress and safety calls where applicable. Entries must be made giving details of all work performed which may affect the proper operation of the station. Logs must be retained by the licensee for a period of two years from the date of entry, and, where applicable, for such additional periods such as logs relating to a distress situation or disaster must be retained for three years from the date of entry in the log. If the Commission has notified the licensee of an investigation, the related logs must be retained until the licensee is specifically authorized in writing to destroy them. Logs relating to any claim or complaint of which the station licensee has notice must be retained until the claim or complaint has been satisfied or barred by statute limiting the time for filing suits upon such claims.

*Section 80.409(d), Ship Radiotelegraph Logs:* Logs of ship stations which are compulsorily equipped for radiotelegraphy and operating in the band 90 to 535 kHz must contain specific information in log entries according to this subsection.

*Section 80.409(e), Ship Radiotelephone Logs:* Logs of ship stations which are compulsorily equipped for radiotelephony must contain specific information in applicable log entries and the time of their occurrence.

The recordkeeping requirements contained in section 80.409 is necessary to document the operation and public correspondence service of public coast radiotelegraph, public coast radiotelephone stations and Alaska-public fixed stations, ship radiotelegraph, ship radiotelephone and applicable radiotelephone including the

logging of distress and safety calls where applicable.

*OMB Control Number:* 3060–0674.

*Title:* Section 76.1618, Basic Tier Availability.

*Type of Review:* Extension of a currently approved collection.

*Respondents:* Business or other for-profit entities.

*Number of Respondents and Responses:* 4,139 respondents; 4,139 responses.

*Estimated Time per Response:* 2.25 hours.

*Frequency of Response:* Third party disclosure requirement.

*Obligation to Respond:* Required to obtain or retain benefits. The statutory authority for this collection of information is contained in Section 4(i) and Section 632 of the Communications Act of 1934, as amended.

*Total Annual Burden:* 9,313 hours.

*Total Annual Cost:* No cost.

*Needs and Uses:* The information collection requirements contained in 47 CFR 76.1618 state that a cable operator shall provide written notification to subscribers of the availability of basic tier service to new subscribers at the time of installation. This notification shall include the following information: (a) That basic tier service is available; (b) the cost per month for basic tier service; and (c) a list of all services included in the basic service tier. These notification requirements are to ensure the subscribers are made aware of the availability of basic cable service at the time of installation.

*OMB Control Number:* 3060–0703.

*Title:* Determining Costs of Regulated Cable Equipment and Installation, FCC Form 1205.

*Form Number:* FCC Form 1205.

*Type of Review:* Revision of a currently approved collection.

*Respondents:* Business or other for-profit entities.

*Number of Respondents and Responses:* 400 respondents; 600 responses.

*Estimated Time per Response:* 4–12 hours.

*Frequency of Response:*

Recordkeeping requirement, Annual reporting requirement, Third party disclosure requirement.

*Obligation to Respond:* Required to obtain or retain benefits. The statutory authority for this collection of information is contained in Section 301(j) of the Telecommunications Act of 1996 and 623(a)(7) of the Communications Act of 1934, as amended.

*Total Annual Burden:* 4,800 hours.

*Total Annual Cost:* \$360,000.

*Needs and Uses:* Information derived from FCC Form 1205 filings is used to

facilitate the review of equipment and installation rates. This information is then reviewed by each cable system's respective local franchising authority. Section 76.923 records are kept by cable operators in order to demonstrate that charges for the sale and lease of equipment for installation have been developed in accordance with the Commission's rules. The Commission modified the language in 47 CFR 76.923 in 90 FR 31145 (July 14, 2025), FCC 25–33, to exclude from regulation equipment that is used to receive tiers other than the basic tier of service. Due to this exclusion, fewer FCC Form 1205 filings will be filed with the Commission. Therefore, Commission is requesting approval of this revision to the information collection that reduces the paperwork burden.

Federal Communications Commission.

**Marlene Dortch,**

*Secretary, Office of the Secretary.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10718, CMS–10611 and CMS–10260]

### 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 24, 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–10718—Model Medicare Advantage and Medicare Prescription Drug Plan Individual Enrollment Request Form**

**CMS–10611—Medicare Outpatient Observation Notice (MOON)**

**CMS–10260—Medicare Advantage and Prescription Drug Program**

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 with of a currently approved collection; **Title of Information Collection:** Model Medicare Advantage and Medicare Prescription Drug Plan Individual Enrollment Request Form; **Use:** The enrollment form is considered a “model” under Medicare regulations at §§ 422.2267 and 423.2267, for purposes of communication and marketing review and approval; therefore, MA and Part D plans are able to modify the language, format, or order of the enrollment form. The model enrollment form includes the minimal amount of information to process the enrollment, located in Section 1 of the MA/PDP enrollment form, and other limited information, in Section 2, that the sponsor is required (i.e., accessible format preference) or chooses (i.e., premium payment information) to provide to the beneficiary. The optional data elements, which aids the MA and Part D plan in processing the enrollment, is developed for efficiency for the plan. Plan sponsors can obtain information at the initial point of contact to help streamline the beneficiary’s enrollment process. The optional questions include information specific to the plan’s business needs that serves to reduce overall burden and allow for timely processing of an enrollment request. All data elements in Section 2 are optional for the beneficiary to complete. Plan enrollment will not be affected if the beneficiary does not complete this additional information. **Form Number:** CMS–10718 (OMB control number 0938–1378); **Frequency:** Occasionally; **Affected Public:** Individuals and Households, Private sector and Business or other for-profits; **Number of Respondents:** 24,464,437; **Number of Responses:** 49,917,959; **Total Annual Hours:** 12,240,174. (For questions regarding this collection, contact: AnhViet Nguyen at (667) 290–9745 or [anhviet.nguyen@cms.hhs.gov](mailto:anhviet.nguyen@cms.hhs.gov).)

2. *Type of Information Collection*  
**Request:** Extension of a currently

approved collection; **Title of Information Collection:** Hospital Notice: Medicare Outpatient Observation Notice (MOON); **Use:** The Medicare Outpatient Observation Notice (MOON), serves as the written notice component of this mandatory notification process. The standardized content of the MOON includes all informational elements required by statute, in language understandable to beneficiaries, and fulfills the regulatory requirements at 42 CFR part 489.20(y).

The MOON is a standardized notice delivered to people entitled to Medicare benefits under Title XVIII of the Act who receive more than 24 hours of observation services, informing them that their hospital stay is outpatient and not inpatient, and the implications of being an outpatient. **Form Number:** CMS–10611 (OMB control number 0938–1308); **Frequency:** Annually; **Affected Public:** Private sector, Business or other for-profits and Not-for-profits institutions; **Number of Respondents:** 5,817; **Number of Responses:** 2,073,991; **Total Annual Hours:** 518,498. (For questions regarding this collection, contact: Katherine Hosna at 410–786–4993 or [Katherine.Hosna@cms.hhs.gov](mailto:Katherine.Hosna@cms.hhs.gov).)

3. *Type of Information Collection*  
**Request:** Revision of a currently approved collection; **Title of Information Collection:** Medicare Advantage and Prescription Drug Program; **Use:** CMS requires MA organizations and Part D sponsors to use the standardized documents being submitted for OMB approval to satisfy disclosure requirements mandated by section 1851 (d)(3)(A) of the Act and § 422.111 for MA organizations and section 1860D–1(c) of the Act and § 423.128(a)(3) for Part D sponsors.

The regulatory provisions at §§ 422.111(b) and 423.128(b) require MA organizations and Part D sponsors to disclose plan information, including: service area, benefits, access, grievance and appeals procedures, and quality improvement/assurance requirements. MA organizations and sponsors may send the ANOC separately from the EOC but must send the ANOC for enrollee receipt by September 30. The required due date for the EOC is 15 days prior to the start of the AEP. **Form Number:** CMS–10260 (OMB control number 0938–1051); **Frequency:** Annually; **Affected Public:** Private sector and Business or other for-profits; **Number of Respondents:** 712; **Number of Responses:** 45,996; **Total Annual Hours:** 12,316. (For questions regarding this collection, contact: Lauren Yeary at

(410) 786-3211 or [lauren.dulay@cms.hhs.gov](mailto:lauren.dulay@cms.hhs.gov)).

**William N. Parham, III,**

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

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**

**[Document Identifier: CMS-10749, CMS-8550, CMS-10328, CMS-10148 and CMS-10572]**

**Agency Information Collection Activities: Submission for OMB Review; 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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by October 24, 2025.

**ADDRESSES:** 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](http://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.

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 Parham at (410) 786-4669.

**SUPPLEMENTARY INFORMATION:** Under the Paperwork Reduction Act of 1995 (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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment.

**1. Type of Information Collection Request:** Reinstatement without change of a previously approved collection; **Title of Information Collection:** National Plan and Provider Enumeration System (NPPES) Supplemental Data Collection; **Use:** The adoption by the Secretary of HHS of the standard unique health identifier for health care providers is a requirement of the Health Insurance Portability and Accountability Act of 1996 (HIPAA). The unique identifier is to be used on standard transactions and may be used for other lawful purposes in the health care system. The CMS Final Rule published on January 23, 2004 adopts the National Provider Identifier (NPI) as the standard unique health identifier for health care providers. Health care providers that are covered entities under HIPAA must apply for and use NPIs in standard transactions. The law requires that data collection standards for these measures be used, to the extent that it is practical, in all national population health surveys. It applies to self-reported optional information only. The law also

requires any data standards published by HHS to comply with standards created by the Office of Management and Budget (OMB).

The web based optional data fields can be seen in Appendix A1: Data Collected for the Office of Minority and Appendix A2: Data collected for the 21st Century Cures Act, interoperability. The standards apply to population health surveys sponsored by HHS, where respondents either self-report information or a knowledgeable person responds for all members of a household. HHS is implementing these data standards in all new surveys. **Form Number:** CMS-10749 (OMB control number: 0938-1427); **Frequency:** Yearly; **Affected Public:** Private Sector, Business or other for-profits, Not-for-profit institutions; **Number of Respondents:** 545,648; **Total Annual Responses:** 545,648; **Total Annual Hours:** 92,760. (For policy questions regarding this collection contact Nora Simmons at 410-786-1981.)

**2. Type of Information Collection Request:** Revision of a currently approved collection; **Title of Information Collection:** Medicare Registration Application; **Use:** Various sections of the Social Security Act (Act), the United States Code (U.S.C.), Internal Revenue Service Code (Code) and the Code of Federal Regulations (CFR) require providers and suppliers to furnish information concerning the amounts due and the identification of individuals or entities that furnish medical services to beneficiaries before allowing payment. The principal function of the CMS-855O is to gather information from a physician or other eligible professional to help CMS determine whether he or she meets certain qualifications to enroll in the Medicare program for the sole purpose of ordering or certifying certain Medicare items or services. The CMS-855O allows a physician or other eligible professional to enroll in Medicare without approval for billing privileges.

The collection and verification of this information protects our beneficiaries from illegitimate providers/suppliers. These procedures also protect the Medicare Trust Funds against fraud. The CMS-855O gathers information that allow Medicare contractors to ensure that the physician or eligible professional is not sanctioned from the Medicare and/or Medicaid program(s), or debarred, or excluded from any other Federal agency or program. Furthermore, the data collected also ensures that the applicant has the necessary credentials to order and certify health care services. This is the