This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at <a href="https://www.federalreserve.gov/foia/request.htm">https://www.federalreserve.gov/foia/request.htm</a>. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)).

Comments received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than October 14, 2025.

A. Federal Reserve Bank of Dallas (Lindsey Wieck, Director, Mergers & Acquisitions) 2200 North Pearl Street, Dallas, Texas 75201–2272. Comments can also be sent electronically to Comments.applications@dal.frb.org:

1. First Federal Community, MHC, Paris, Texas; to become a bank holding company by acquiring First Federal Community Bank, SSB, Paris Texas, upon the conversion of First Federal Community Bank, SSB, from mutual to stock form.

Board of Governors of the Federal Reserve System.

#### Michele Taylor Fennell,

Associate Secretary of the Board. [FR Doc. 2025–17654 Filed 9–11–25; 8:45 am] BILLING CODE P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10923 and CMS-116]

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 14, 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. 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: New collection (Request for a new OMB control number); Title: Generic Clearance for Improving DASG's APIs Customer Experience; Use: This new request will enable the Centers for Medicare & Medicaid Services' Data Analytics and Systems Group (hereafter "the Agency") to act in accordance with OMB Circular A-11 Section 280, Executive Order 14058, and the OMB memoranda on burden reduction, to ultimately transform the experience of its customers to improve both efficiency and mission delivery and increase accountability by communicating about these efforts with the public.

The Agency will collect, analyze, and interpret information gathered through this generic clearance to identify services' accessibility, navigation, and use by customers and make improvements in service delivery based on customer insights gathered through developing an understanding of the user experience interacting with the Government. Under this request, two types of activities (Customer Research and Customer Feedback Surveys) will be conducted to generate customer insights. Form Number: CMS-10923 (OMB control number: 0938-New): Frequency: Occasionally; Affected Public: Individual and Households, Private sector (Business or other forprofits), Not for profits institutions; Number of Respondents: 82,020; Number of Responses: 82,020; Total Annual Hours: 15,876. (For questions regarding this collection, contact Rachel Lorencz at rachel.lorencz@cms.hhs.gov.

2. Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: Clinical Laboratory Improvement Amendments (CLIA) Application Form and Supporting Regulations; *Use:* Section 353(b) of the Public Health Service Act specifies that the laboratory must submit an application in such form and manner as the Secretary shall prescribe that describes the characteristics of the laboratory and examinations and procedures performed by the laboratory. The application must be completed by entities performing laboratory's testing specimens for diagnostic or treatment

purposes. This information is vital to the certification process. In this revision, changes were made to the form to facilitate its completion and data entry. We anticipate that the changes will not increase the time to complete the form. Form Number: CMS-116 (OMB control number: 0938-0581); Frequency: Biennially and Occasionally; Affected Public: Private Sector-Business or other for-profits and Notfor-profit institutions; Number of Respondents: 50,236; Total Annual Responses: 50,236; Total Annual Hours: 50,236. (For policy questions regarding this collection contact Cheryl Murphy at Cheryl.Murphy@cms.hhs.gov.)

### William N. Parham, III,

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

[FR Doc. 2025-17660 Filed 9-11-25; 8:45 am]

BILLING CODE 4120-01-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10065/10066 and CMS-10690]

### 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, 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 <a href="http://www.regulations.gov">http://www.regulations.gov</a>. 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

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-10065/10066 Hospital Notices: IM/DND

CMS-10690 CLIA Proficiency Testing (PT)

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: Extension of a currently approved collection; Title of Information Collection: Hospital Notices: IM/DND; Use: This information collection applies to beneficiaries in Original Medicare and enrollees in Medicare health plans. The purpose of the IM is to inform beneficiaries and enrollees of their rights as hospital inpatients and how to request a discharge appeal by a Quality Improvement Organization (QIO) and how to file a request. Consistent with 42 CFR 405.1205 for Original Medicare and 422.620 for Medicare health plans, hospitals must provide the initial IM within 2 calendar days of admission. A follow-up copy of the signed IM is given no more than 2 calendar days before discharge. The follow-up copy is not required if the first IM is provided within 2 calendar days of discharge. Form Number: CMS-10065/10066 (OMB control number: 0938-1019); Frequency: Yearly; Affected Public: Private Sector, Business or other for profits, Not for profit institutions; Number of Respondents: 25,397,156; Total Annual Responses: 25,397,156; Total Annual Hours: 4,313,823. (For policy questions regarding this collection contact: Katherine Hosna at 410-786-4993 or KatherineHosna@ cms.hhs.gov).

2. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: CLIA Proficiency Testing (PT); Use: This is an extension package. The purpose of this package is to request Office of Management and Budget (OMB) approval for the information collection request (ICR) for proficiency testing (PT) and reapproval of PT programs. The ICR includes laboratories filling in PT submission forms for microbiology PT and document collection for a PT program if it needs to reapply for approval using the initial approval process.

On October 31, 1988, Congress enacted the Clinical Laboratory Improvement Amendments of 1988 (Pub. L. 100–578) (CLIA'88), codified at 42 U.S.C. 263a, to ensure the accuracy and reliability of testing in all laboratories, including, but not limited to, those that participate in Medicare