Observers requiring auxiliary aids should email *DisabilityProgram@fdic.gov* to make necessary arrangements.

STATUS: Open to public observation via webcast.

MATTERS TO BE CONSIDERED: The Federal Deposit Insurance Corporation's (FDIC) Board of Directors will meet to consider the following matters:

Discussion Agenda

Notice of Proposed Rulemaking: Regulatory Capital Rule: Revisions to the Community Bank Leverage Ratio Framework.

Final Rule: Regulatory Capital Rule: Modifications to the Enhanced Supplementary Leverage Ratio Standards for U.S. Global Systemically Important Bank Holding Companies and Their Subsidiary Depository Institutions; Total Loss-Absorbing Capacity and Long-Term Debt Requirements for U.S. Global Systemically Important Bank Holding Companies.

Summary Agenda

No substantive discussion of the following items is anticipated. The Board of Directors will resolve these matters with a single vote unless a member of the Board requests that an item be moved to the discussion agenda.

Final Rule: Adjusting and Indexing Certain Regulatory Thresholds.

Designated Reserve Ratio for 2026.

Final Rule; Delay of Compliance Date: FDIC Official Signs and Advertising Requirements, False Advertising, Misrepresentation of Insured Status, and Misuse of the FDIC's Name or Logo.

Minutes of a Board of Directors' Meeting Previously Distributed.

Summary reports, status reports, and reports of actions taken pursuant to authority delegated by the Board of Directors.

CONTACT PERSON FOR MORE INFORMATION: For further information, please contact Debra A. Decker, Executive Secretary, FDIC, at *FDICBoardMatters@fdic.gov.*

Authority: 5 U.S.C. 552b.

Dated at Washington, DC, on November 19, 2025.

Federal Deposit Insurance Corporation.

Debra A. Decker,

Executive Secretary.

[FR Doc. 2025–20728 Filed 11–20–25; 11:15 am]

BILLING CODE 6714–01–P

FEDERAL TRADE COMMISSION

SES Performance Review Board

AGENCY: Federal Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given of the appointment of members to the FTC Performance Review Board.

FOR FURTHER INFORMATION CONTACT: Tamika Williams, Chief Human Capital Officer, 600 Pennsylvania Avenue NW, Washington, DC 20580, (202) 326–2184.

SUPPLEMENTARY INFORMATION: Publication of the Performance Review Board (PRB) membership is required by 5 U.S.C. 4314(c)(4). The PRB reviews and evaluates the initial appraisal of a senior executive's performance by the supervisor, and makes recommendations regarding performance ratings, performance awards, and pay-for-performance pay adjustments to the Chair.

The following individuals have been designated to serve on the Commission's Performance Review Board:

Lucas Croslow, General Counsel, PRB Chair
Daniel Guarnera, Director, Bureau of Competition
Christopher Mufarrige, Director, Bureau of Consumer Protection
David Robbins, Executive Director
Ted Rosenbaum, Deputy Director for Research and Management, Bureau of Economics
Rebecca Unruh, Deputy Director, Bureau of Consumer Protection
Katherine White, Deputy Director, Bureau of Consumer Protection
Tamika Williams, Chief Human Capital Officer

By direction of the Commission.

Joel Christie,

Acting Secretary.

[FR Doc. 2025–20761 Filed 11–21–25; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10781]

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 December 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.* 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:* FOIA/Privacy Act Requests for Medicare Claims Data via CMS FOIA Public Portal; *Use:* This collection of information is dedicated to Medicare beneficiaries and third-party requesters (law firms or others) acting on behalf of beneficiaries that are making requests for CMS to produce Medicare beneficiary records through 5 U.S.C. 552(b) (See also 42 CFR 401.136). The online portal allows for ease and efficiency in uploading requests and required authorizations. Additionally, with the portal, requesters can securely submit requests electronically that contain PHI or PII. They are advised that *MyMedicare.gov/Blue Button3* is an online service available for beneficiaries to set up an account to access their own records and give authorization to share with third parties. This secure public online portal is integrated with CMS's current FOIA/Privacy Act case management system to enter, track, and process incoming FOIA requests (See 45 CFR 5.22 and 5.24). Unless permitted or required by law, the Health Insurance Portability and Accountability Act (HIPAA) Privacy Rule (45 CFR 164.508) prohibits Medicare (a HIPAA-covered entity) from disclosing an individual's protected health information without valid authorization. *Form Number:* CMS–10781 (OMB control number: 0938–1419); *Frequency:* Reporting—Occasionally; *Affected Public:* Individuals or Households; *Number of Respondents:* 22,600; *Total Annual Responses:* 22,600; *Total Annual Hours:* 7,533. (For policy questions regarding this collection contact Joseph Tripline at *joseph.tripline@cms.hhs.gov*).

William N. Parham, III,

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

[FR Doc. 2025–20705 Filed 11–21–25; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10849]

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 December 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.* 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:* Revision of a currently approved collection; *Title of Information Collection:* Drug Price Negotiation for Initial Price Applicability Year 2028 under Sections 11001 and 11002 of the Inflation Reduction Act; *Use:* Under the authority in sections 11001 and 11002 of the Inflation Reduction Act of 2022 (*Pub. L. 117–169*), the Centers for Medicare & Medicaid Services (CMS) is implementing the Medicare Drug Price Negotiation Program, codified in sections 1191 through 1198 of the Social Security Act (''the Act''). The Act establishes the Negotiation Program to negotiate maximum fair prices (''MFPs''), defined at 1191(c)(3) of the Act, for certain high expenditure, single source selected drugs covered under Medicare Part B and Part D. For the third cycle of the Negotiation Program, the Secretary of Health and Human Services (the ''Secretary'') will select up to 15 high expenditure, single source drugs payable under Part B and/or covered under Part D for negotiation. In accordance with section 1194(f)(4) of the Act, CMS will also renegotiate MFPs for drugs selected for renegotiation, if any, for initial price applicability year 2028.

Negotiation Data Elements: The statute requires that CMS consider certain data from Primary Manufacturers as part of the negotiation process. To the extent that more than one entity meets the statutory definition of manufacturer (specified in section