the information has practical utility; (b) the accuracy of the estimates of the burden of the information collection, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. All comments will become a matter of public record.

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on August 21, 2025.

Jennifer M. Jones,

Deputy Executive Secretary.

[FR Doc. 2025–16200 Filed 8–22–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10434 #26 and #47]

Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit 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 September 8, 2025.

ADDRESSES: When commenting, please reference the applicable form number (CMS–10434 #____) and the OMB control number (0938–1188). 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: CMS–10434____/OMB control number: 0938–1188, Room C4–26–05, 7500 Security Boulevard, Baltimore, MD 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/medicare/regulations-guidance/legislation/paperwork-reduction-act-1995/pra-listing*.

FOR FURTHER INFORMATION CONTACT: William N. Parham at 410–786–4669.

SUPPLEMENTARY INFORMATION: Following is a summary of the use and burden associated with the subject information collection(s). More detailed information can be found in the collection's supporting statement and associated materials (see **ADDRESSES**).

Generic Information Collection

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Child and Adult Core Set Measures; *Use:* CMS is required annually to revise and update the Child and Adult Core Sets. The review for the 2025 Core Set resulted in several substantive changes to the templates used for state Core Set reporting to CMS. State reporting of the Core Sets in the Quality Measurement Reporting system opens for 2025 Core Set reporting on September 3, 2025. *Form Number:* CMS–10434 #26 (OMB control number: 0938–1188); *Frequency:* Yearly, once, and occasionally; *Affected Public:* Individuals or households and State, Local, or Tribal Governments; *Number of Respondents:* 61,293; *Total Annual Responses:* 61,455; *Total Annual Hours:* 99,977. (For policy questions regarding this collection contact: Virginia (Gigi) Raney at 410–786–6117.)

2. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Health Home Core Sets Measures; *Use:* CMS is required annually to revise and update the Health Home Core Sets. The review for the 2025 Core Set resulted in several substantive changes to the templates used for state Core Set reporting to CMS. State reporting of the Core Sets in the Quality Measurement Reporting system opens for 2025 Core Set reporting on September 3, 2025. *Form Number:* CMS–10434 #47 (OMB control number: 0938–1188); *Frequency:* Yearly, once, and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 50; *Total Annual Responses:* 50; *Total Annual Hours:* 4,620. (For policy questions regarding this collection contact: Sara Rhoades at 410–786–4484.)

Evell Barco Holland,

Senior Technical Advisor, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2025–16171 Filed 8–22–25; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–N–2962]

Prescription Drug User Fee Act and Biosimilar User Fee Amendments Hiring and Retention Assessment; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the following public meeting entitled ''Prescription Drug User Fee Act and Biosimilar User Fee Amendments Hiring and Retention Assessment.'' The topic to be discussed is a hiring and retention assessment