appropriately to complaints against accredited facilities.

++ CHAP's processes and procedures for monitoring a HIT supplier found out of compliance with CHAP's program requirements.

++ CHAP's capacity to report deficiencies to the surveyed facilities and respond to the facility's plan of correction in a timely manner.

++ CHAP's capacity to provide CMS with electronic data and reports necessary for effective assessment and interpretation of the organization's survey process.

++ The adequacy of CHAP's staff and other resources, and its financial viability.

++ CHAP's capacity to adequately fund required surveys.

++ CHAP's policies with respect to whether surveys are announced or unannounced, to ensure that surveys are unannounced.

++ CHAP's agreement to provide CMS with a copy of the most current accreditation survey together with any other information related to the survey as CMS may require (including corrective action plans).

++ CHAP's policies and procedures to avoid conflicts of interest, including the appearance of conflicts of interest, involving individuals who conduct surveys, audits or participate in accreditation decisions.

++ CHAP's agreement or policies for voluntary and involuntary termination of HIT suppliers.

++ CHAP's agreement or policies for voluntary and involuntary termination of the HIT AO program.

IV. Collection of Information Requirements

This document does not impose information collection requirements, that is, reporting, recordkeeping, or third-party disclosure requirements. Consequently, there is no need for review by the Office of Management and Budget under the authority of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*).

V. Response to Comments

Because of the large number of public comments, we normally receive on **Federal Register** documents, we are not able to acknowledge or respond to them individually. We will consider all comments we receive by the date and time specified in the **DATES** section of this preamble, and, when we proceed with a subsequent document, we will respond to the comments in the preamble to that document.

The Administrator of the Centers for Medicare & Medicaid Services (CMS), Chiquita Brooks-LaSure, having reviewed and approved this document, authorizes Trenesha Fultz-Mimms, who is the Federal Register Liaison, to electronically sign this document for purposes of publication in the **Federal Register**.

Trenesha Fultz-Mimms,

Federal Register Liaison, Centers for Medicare & Medicaid Services.

[FR Doc. 2024–09176 Filed 4–29–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10788]

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 *May 30, 2024.*

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:* Prescription Drug and Health Care Spending; *Use:* On December 27, 2020, the Consolidated Appropriations Act, 2021 (CAA) was signed into law. Section 204 of Title II of Division BB of the CAA added parallel provisions at section 9825 of the Internal Revenue Code (the Code), section 725 of the Employee Retirement Income Security Act (ERISA), and section 2799A–10 of the Public Health Service Act (PHS Act) that require group health plans and health insurance issuers offering group or individual health insurance coverage to annually report to the Department of the Treasury, the Department of Labor (DOL), and the Department of Health and Human Services (HHS) (collectively, ''the Departments'') certain information about prescription drug and health care spending, premiums, and enrollment under the plan or coverage. This information will support the

development of public reports that will be published by the Departments on prescription drug reimbursements for plans and coverage, prescription drug pricing trends, and the role of prescription drug costs in contributing to premium increases or decreases under the plans or coverage. The 2021 interim final rules, ''Prescription Drug and Health Care Spending'' (2021 interim final rules), issued by the Departments and the Office of Personnel Management (OPM) implement the provisions of section 9825 of the Code, section 725 of ERISA, and section 2799A–10 of the PHS Act, as enacted by section 204 of Title II of Division BB of the CAA. OPM joined the Departments in issuing the 2021 interim final rules, requiring Federal Employees Health Benefits (FEHB) carriers to report information about prescription drug and health care spending, premiums, and plan enrollment in the same manner as a group health plan or health insurance issuer offering group or individual health insurance coverage.

The 2023 Prescription Drug Data Collection (RxDC) Reporting Instructions reflect changes for the 2023 reference year and beyond. As a result of removing first-year implementation costs and burdens that were incurred prior to 2024, it is estimated that there will be a decrease in total three-year average annual burden from 1,684,080 to 668,952. *Form Number:* CMS–10788 (OMB Control Number: 0938–1407); *Frequency:* Annually; *Affected Public:* Private Sector; *Number of Respondents:* 356; *Number of Responses:* 356; *Total Annual Hours:* 668,952. (For policy questions regarding this collection contact Christina Whitefield at 202–536–8676.)

William N. Parham, III,

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

[FR Doc. 2024–09314 Filed 4–29–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for Office of Management and Budget (OMB) Review; Generic Clearance for the Comprehensive Child Welfare Information System (CCWIS) Technical Assistance and Review Process (OMB #: 0970–0568)

AGENCY: Children's Bureau, Administration for Children and Families, U.S. Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Children's Bureau (CB), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS) is requesting a 3-year extension of the Generic Clearance for the Comprehensive Child Welfare Information System (CCWIS) Technical Assistance (TA) and Review Process, (OMB #0970–0568, expiration 4/30/2024) and all approved information collections under this generic. There are no changes requested to the terms of the umbrella generic or to the currently approved information collections.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

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. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The CCWIS Technical Assistance and Review information collection includes two components.

- The CCWIS Assessment Review (CAR) Process.
- TA tools for title IV–E agencies to self-assess their conformity to CCWIS project and design requirements at 45 CFR 1355.52–3.

The CCWIS requirements at 45 CFR 1355.55 require the review, assessment, and inspection of the planning, design, development, installation, operation, and maintenance of each CCWIS project on a continuing basis. The Advance Planning Document (APD) regulations at 45 CFR 95.621 require periodic reviews of state and local agency methods and practices to ensure information systems, including CCWIS, are utilized for purposes consistent with proper and efficient administration.

This request is for an extension with no changes to the umbrella generic and all currently approved information collections, which can be found here: *https://www.reginfo.gov/public/do/PRAICList?ref_nbr=202311-0970-010.*

Respondents: Title IV–E agencies under the Social Security Act.

Annual Burden Estimates

ANNUAL BURDEN—CURRENTLY APPROVED INFORMATION COLLECTIONS

Instrument	Total number of respondents	Total number of responses per respondent (3 years)	Average burden hours per response	Total burden hours	Annual burden hours
CCWIS Self-Assessment—Administration	55	1	10	550	183
CCWIS Self-Assessment—Adoption	55	1	10	550	183
CCWIS Self-Assessment—Case Management	55	1	10	550	183
CCWIS Self-Assessment—Foster Care and Service Provider Management	55	1	10	550	183
CCWIS Self-Assessment—Intake	55	1	10	550	183
CCWIS Self-Assessment—Investigation	55	1	10	550	183
CCWIS Self-Assessment: Child Welfare Contributing Agency (CWCA)	55	1	10	550	183
CCWIS Self-Assessment: Data Exchanges	55	1	10	550	183
CCWIS Self-Assessment: Data Quality	55	1	10	550	183