

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Type of data collection instrument	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Intermediary or end users (e.g., employers, workers, manufactures, labor/professional associations, policymakers).	Interview or focus group guide .....	25	1	1	25
	Survey instrument (pre and post) ....	10,000	1	20/60	3,333
	Informed consent form .....	650	1	5/60	54
	Interview or focus group guide .....	650	1	1	650
Total .....	.....	.....	.....	.....	6,069

**Jeffrey M. Zirger,**  
*Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.*  
 [FR Doc. 2024-14309 Filed 6-27-24; 8:45 am]  
**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**Advisory Committee on Breast Cancer in Young Women; Notice of Charter Renewal**

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice of charter renewal.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), within the Department of Health and Human Services (HHS), announces the renewal of the charter of the Advisory Committee on Breast Cancer in Young Women (ACBCYW).

**FOR FURTHER INFORMATION CONTACT:** Kimberly E. Smith, M.B.A., M.H.A., Designated Federal Officer, Advisory Committee on Breast Cancer in Young Women, Centers for Disease Control and Prevention, Department of Health and Human Services, 4770 Buford Highway NE, Mailstop S107-4, Atlanta, Georgia 30341-3717. Telephone: (404) 498-0073; Email: [KESmith@cdc.gov](mailto:KESmith@cdc.gov).

**SUPPLEMENTARY INFORMATION:** CDC is providing notice under 5 U.S.C. 1001-1014 of the renewal of the charter of the Advisory Committee on Breast Cancer in Young Women, Centers for Disease Control and Prevention, Department of Health and Human Services. This charter has been renewed for a two-year period through June 17, 2026.

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been

delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**  
*Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.*  
 [FR Doc. 2024-14289 Filed 6-27-24; 8:45 am]

**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**

**[Document Identifiers: CMS-179, CMS-10536, CMS-R-153 and CMS-10326]**

**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 August 27, 2024.

**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-179 Medicaid State Plan Base Plan Pages

CMS-10536 Medicaid Eligibility and Enrollment (EE) Implementation Advanced Planning Document (IAPD) Template

CMS-R-153 Medicaid Drug Use Review (DUR) Program

CMS-10326 Electronic Submission of Medicare Graduate Medical Education (GME)

### Affiliation Agreements

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:* Medicaid State Plan Base Plan Pages; *Use:* State Medicaid agencies complete the plan pages while we review the information to determine if the state has met all of the requirements of the provisions the states choose to implement. If the requirements are met, we will approve the amendments to the state's Medicaid plan giving the state the authority to implement the flexibilities. For a state to receive Medicaid Title XIX funding, there must be an approved Title XIX state plan. *Form Number:* CMS-179 (OMB control number 0938-0193); *Frequency:* Occasionally; *Affected Public:* State, Local, and Tribal Governments; *Number of Respondents:* 56; *Total Annual Responses:* 1,120; *Total Annual Hours:* 22,400. (For policy questions regarding this collection contact Gary Knight at 304-347-5723.)

#### 2. Type of Information Collection

*Request:* Extension of a currently approved collection; *Title of Information Collection:* Medicaid

Eligibility and Enrollment (EE) Implementation Advanced Planning Document (IAPD) Template; *Use:* To assess the appropriateness of states' requests for enhanced federal financial participation for expenditures related to Medicaid eligibility determination systems, we will review the submitted information and documentation to make an approval determination for the advanced planning document. *Form Number:* CMS-10536 (OMB control number: 0938-1268); *Frequency:* Yearly, once, and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 56; *Total Annual Responses:* 168; *Total Annual Hours:* 2,688. (For policy questions regarding this collection contact Loren Palestino at 410-786-8842.)

#### 3. Type of Information Collection

*Request:* Revision of a currently approved collection; *Title of Information Collection:* Medicaid Drug Use Review (DUR) Program; *Use:* States must provide for a review of drug therapy before each prescription is filled or delivered to a Medicaid patient. This review includes screening for potential drug therapy problems due to therapeutic duplication, drug-disease contraindications, drug-drug interactions, incorrect drug dosage or duration of drug treatment, drug-allergy interactions, and clinical abuse/misuse. Pharmacists must make a reasonable effort to obtain, record, and maintain Medicaid patient profiles. These profiles must reflect at least the patient's name, address, telephone number, date of birth/age, gender, history, e.g., allergies, drug reactions, list of medications, and pharmacist's comments relevant to the individual's drug therapy. The State must conduct retrospective drug use review which provides for the ongoing periodic examination of claims data and other records in order to identify patterns of fraud, abuse, inappropriate or medically unnecessary care. Patterns or trends of drug therapy problems are identified and reviewed to determine the need for intervention activity with pharmacists and/or physicians. States may conduct interventions via telephone, correspondence, or face-to-face contact. The states and managed care organizations (MCOs) are provided the reporting instrument (a survey) by CMS, and by responding to the survey, the states generate annual reports which are submitted to CMS for the purposes of monitoring compliance and evaluating the progress of states' DUR programs. The survey and the annual recordkeeping and reporting requirements under the pertinent regulations, are completed by

pharmacists employed by, or contracted with the various state Medicaid programs and their MCOs. The annual reports submitted by states are reviewed and results are compiled by CMS in a format intended to provide information, comparisons and trends related to states' experiences with DUR. The states benefit from the information and may enhance their programs each year based on state reported innovative practices that are compiled by CMS from the annual reports. A comparison/summary of the data from the annual reports is published on Medicaid.gov annually, and serves as a resource for stakeholders, including but not limited to states, manufacturers, researchers, congress, CMS, the Office of Inspector General, non-governmental payers and clinicians on the topic of DUR in state Medicaid programs. *Form Number:* CMS-R-153 (OMB control number: 0938-0659); *Frequency:* Yearly, quarterly, and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 52; *Total Annual Responses:* 676; *Total Annual Hours:* 41,860. (For policy questions regarding this collection contact Mike Forman at 410-786-2666.)

#### 4. Type of Information Collection

*Request:* Reinstatement without change of a currently approved collection; *Title of Information Collection:* Electronic Submission of Medicare Graduate Medical Education (GME) Affiliation Agreements; *Use:* Existing regulations at § 413.75(b) permit hospitals that share residents to elect to form a Medicare GME affiliated group if they are in the same or contiguous urban or rural areas, if they are under common ownership, or if they are jointly listed as program sponsors or major participating institutions in the same program by the accrediting agency. The purpose of a Medicare GME affiliated group is to provide flexibility to hospitals in structuring rotations under an aggregate full time equivalent (FTE) resident cap when they share residents. The existing regulations at § 413.79(f)(1) specify that each hospital in a Medicare GME affiliated group must submit a Medicare GME affiliation agreement (as defined under § 413.75(b)) to the Medicare Administrative Contractor (MAC) servicing the hospital and send a copy to the Centers for Medicare and Medicaid Services' (CMS) Central Office, no later than July 1 of the residency program year during which the Medicare GME affiliation agreement will be in effect.

CMS will use the information contained in electronic affiliation agreements as documentation of the existence of Medicare GME affiliations,

and to verify that the affiliations being formed by teaching hospitals for the purposes of sharing their Medicare GME FTE cap slots are valid according to CMS regulations. CMS will also use these affiliation agreements as reference materials when potential issues involving specific affiliations arise. While we have used hard copies of affiliation agreements for those same purposes in the past, we implemented this electronic submission process in order to expedite and ease the process of retrieving, analyzing and evaluating affiliation agreements. *Form Number:* CMS-10326 (OMB control number: 0938-1111); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for profits, Not for profit institutions; *Number of Respondents:* 125; *Total Annual Responses:* 125; *Total Annual Hours:* 166. (For policy questions regarding this collection contact Shevi Marciano at 410-786-2874.)

**William N. Parham, III,**

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

[FR Doc. 2024-14338 Filed 6-27-24; 8:45 am]

**BILLING CODE 4120-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for Office of Management and Budget Review; Refugee Support Services (RSS) and RSS Set Aside Sub-Agency List (Office of Management and Budget #0970-0556)**

**AGENCY:** Office of Refugee Resettlement, Administration for Children and Families, U.S. Department of Health and Human Services.

**ACTION:** Request for public comments.

**SUMMARY:** The Administration for Children and Families (ACF) Office of Refugee Resettlement (ORR) is requesting a 3-year extension of the Refugee Support Services (RSS) and RSS Set Aside Sub-Agency List (Office of Management and Budget #0970-0556). ORR is not proposing any changes to the form.

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

**SUPPLEMENTARY INFORMATION:**

*Description:* The RSS and RSS Set Aside Sub-Agency List requests grantees to provide the agency name, city, state, website, and funding amount for each contracted sub-grantee.

The information will be used for national resource mapping pertaining to ORR RSS funding at the local level. Improved communication and the knowledge of all local providers is important to ORR’s overall oversight of the program. In addition to RSS formula funding to states and state replacement agencies who then issue sub-awards to local providers, ORR also awards discretionary grants that directly fund local refugee service providers. This report will continue to provide ORR a complete picture of the availability all ORR resources to assist newly arrived refugees at the local level increasing our ability to identify gaps or target areas of need.

*Respondents:* State agencies and replacement designees under 45 CFR 400.301(c) administering or supervising the administration of programs.

**ANNUAL BURDEN ESTIMATES**

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
RSS and RSS Set Aside Sub-grantee List .....	59	1	2	118

*Authority:* Refugee Act of 1980 [Immigration and Nationality Act, title IV, chapter 2, section 412 (e)] and 45 CFR 400.28.

**Mary C. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2024-14165 Filed 6-27-24; 8:45 am]

**BILLING CODE 4184-45-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for Office of Management and Budget (OMB) Review; Community Services Block Grant (CSBG) Annual Progress Report (OMB No. 0970-0492)**

**AGENCY:** Office of Community Services, Administration for Children and Families, U.S. Department of Health and Human Services.

**ACTION:** Request for public comments.

**SUMMARY:** The Office of Community Services (OCS), Administration for Children and Families (ACF) requests

an extension with minor changes to the currently approved Community Services Block Grant (CSBG) Annual Progress Report, (OMB #0970-0492, expiration 6/30/2024) and is submitting the Tribal Annual Report and Tribal Short Form, as well as removing supplemental funding reports that are no longer in use. Plans for further revisions to this report are also discussed below.

**DATES:** *Comments due* July 29, 2024.

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.