this document for purposes of publication in the **Federal Register**.

Chyana Woodyard,

Federal Register Liaison, Centers for Medicare & Medicaid Services.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10704]

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 July 14, 2026.

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**).

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 Collection

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Health Reimbursement Arrangements and Other Account-Based Group Health Plans; *Use:* On June 20, 2019, the Department of the Treasury, the Department of Labor, and the Department of Health and Human Services (HHS) (collectively, the Departments) issued final regulations, titled ''Health Reimbursement Arrangements and Other Account-Based Group Health Plans'' (84 FR 28888) (2019 final regulations) under section 2711 of the PHS Act and the health nondiscrimination provisions of HIPAA, Public Law 104–191 (HIPAA nondiscrimination provisions). The 2019 final regulations expanded the use of health reimbursement arrangements and other account-based group health plans (collectively referred to as HRAs) and recognized certain HRAs as limited excepted benefits (the excepted benefit HRA), for plan years beginning on or after January 1, 2020. In general, the 2019 final regulations expanded the use of HRAs by eliminating the prohibition on integrating HRAs with individual health insurance coverage, thereby permitting employers to offer individual coverage HRAs to employees that can be integrated with individual health insurance coverage or Medicare Parts A and B, or Part C. Under the 2019 final regulations, employees are permitted to use amounts in an individual coverage HRA to pay expenses for medical care (including premiums for individual health insurance coverage and Medicare), subject to certain requirements.

The information collections associated with the 2019 final regulations are related to the substantiation requirements for individual coverage HRAs (45 CFR 146.123(c)(5)), the notice requirement for individual coverage HRAs (45 CFR 146.123(c)(6)), and notification of termination of coverage (45 CFR 146.123(c)(1)(iii)). Under final regulations issued by HHS on May 14, 2020, titled ''Patient Protection and Affordable Care Act; HHS Notice of Benefit and Payment Parameters for 2021; Notice Requirement for Non-federal Governmental Plans'' (85 FR 29164), under 45 CFR 146.145(b)(3)(viii)(E), excepted benefit HRAs offered by non-Federal governmental plan sponsors are required to provide a notice that describes conditions pertaining to eligibility to receive benefits, annual or lifetime caps or other limits on benefits under the excepted benefit HRA, and a description or summary of the benefits. This notice must be provided no later than 90 days after the employee becomes a participant in the excepted benefit HRA and annually thereafter. *Form Number:* CMS–10704 (OMB Control Number 0938–1361); *Frequency:* Annually; *Affected Public:* State, Local or Tribal governments; *Number of Respondents:* 6,354; *Total Annual Responses:* 1,347,048; *Total Annual Hours:* 3,136. (For policy questions

regarding this collection contact Adam Pellillo at (667) 290–9621.)

William N. Parham, III,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget #: 0970–0473]

Submission for Office of Management and Budget Review; Child Care and Development Fund (CCDF) Consumer Education Website and Reports of Serious Injuries and Death

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

ACTION: Request for public comments.

SUMMARY: The Office of Child Care, Administration for Children and Families (ACF), U.S. Department of Health and Human Services, is requesting a 3-year extension of the Child Care and Development Fund (CCDF) Consumer Education website and Reports of Serious Injuries and Death (Office of Management and Budget)#: 0970–0473, expiration date: May 31, 2026). There are no changes requested to the reporting requirements. Burden estimates have been adjusted.

DATES: *Comments due* June 15, 2026.

ADDRESSES: The public may view and comment on this information collection request at: *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202605-0970-003.* 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 existing *Consumer Education Website reporting requirement* will not be modified and requires states and territories to include information about state or territory policies (related to licensing, monitoring, and background checks) and provider-specific information, including results of monitoring and inspection reports, and if available, information about quality. The existing *Reporting of Serious Injuries and Death reporting requirement* will not be modified. CCDF lead agencies must establish procedures that require child care providers that care for children receiving CCDF subsidies to report to a designated state, territorial, or tribal entity any serious injuries or deaths of children occurring in child care. There are no standard federal forms associated with these reporting requirements.

Respondents: The *Consumer Education Website* requirement applies to the 50 states, the District of Columbia, and 5 territories that receive CCDF grants. *Reporting of Serious Injuries and Death* is a requirement for child care providers receiving CCDF subsidies within states, territories, and tribes.

Annual Burden Estimates

The *burden estimates for the consumer education website* requirement at § 98.33 have been reduced based on analysis of the federal fiscal year 2025–2027 CCDF State and Territory Plans. Eighteen out of 56 lead agencies are non-compliant on requirements related to their consumer education websites and would incur burden under this renewal, while the remaining grantees have demonstrated compliance and would experience minimal burden for ongoing maintenance to their consumer education websites. Accordingly, the burden hours for those 38 lead agencies only needing ongoing maintenance has been reduced from 300 to 50 average hours, resulting in an overall reduction of approximately 57 percent in total annual burden hours compared to the previously approved version. The *burden estimates for reporting serious injuries and deaths* at § 98.42 remain unchanged, as the burden is limited to provider reporting when a reportable event occurs. States and territories have flexibility in how they structure the provider reporting.

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Consumer Education Website Development and Maintenance	18	1	300	5,400
Consumer Education Website Maintenance	38	1	50	1,900
Reporting of Serious Injuries and Death	10,000	1	1	10,000
Estimated Total Annual Burden Hours				17,300

(Authority: Pub. L. 113–186; 42 U.S.C. 9858 *et seq.*)

Mary C. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–N–0421]

Oscar Bobo: Final Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is issuing an order under the Federal Food, Drug, and Cosmetic Act (FD&C Act) debarring Oscar Bobo for a period of 5 years from importing or offering for import any drug into the United States. FDA bases this order on a finding that Mr. Bobo engaged in a pattern of importing or offering for import misbranded drugs (*i.e.,* in an amount, frequency, or dosage that is inconsistent with his personal or household use) that are not designated in an authorized electronic data interchange system as products regulated by FDA. Mr. Bobo was given notice of the proposed debarment and was given an opportunity to request a hearing to show why he should not be debarred. As of March 25, 2026 (30 days after receipt of the notice), Mr. Bobo had not responded. Mr. Bobo's failure to