

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10902]

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 November 4, 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–10902 Environmental Health Hazards Checklist Medicare Coverage for Individuals Exposed to Environmental Health Hazards

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:* New Collection (Request for a new OMB control number); *Title of Information Collection:* Environmental Health Hazards Checklist Medicare Coverage for Individuals Exposed to Environmental Health Hazards; *Use:* Section 1881A of the Act provides an enrollment basis for individuals who have been exposed to environmental health hazards. Currently, the only individuals eligible for Medicare under this provision are those who were present in Lincoln County, Montana, and have an asbestos-related disease (ARD) diagnosis. Eligible individuals must be diagnosed with one or more asbestos-related conditions and have

been present in Lincoln County, Montana, for a total of at least 6 months (need not be consecutive) in the period ending 10 years or more before diagnosis of an asbestos-related condition. This form provides verification from a provider so that SSA can determine eligibility for Medicare enrollment.

SSA uses this information to determine whether an individual meets the requirements for Medicare enrollment on the basis of an Environmental Health Hazard. The form is faxed to the applicant's provider by SSA. The provider must complete and sign the form and submit it back to SSA via fax or mail. The information on the completed form is reviewed manually by SSA. Thus, the collection of this information does not involve the use of information technology. *Form Number:* CMS–10902 (OMB control number: 0938–New); *Frequency:* Once; *Affected Public:* Individuals and Households; *Number of Respondents:* 61; *Total Annual Responses:* 61; *Total Annual Hours:* 10. (For policy questions regarding this collection contact Tyrissa Woods at 410–786–0286 or Tyrissa.Woods@cms.hhs.gov).

William N. Parham, III,

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

[FR Doc. 2024–19867 Filed 9–4–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Child Care Development Fund (CCDF) ACF–696T Financial Report for Tribes (OMB #0970–0510)

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 Administration for Children and Families (ACF) Office of Child Care plans to submit revisions to an approved generic information collection (GenIC) under the umbrella generic: Generic Clearance for Financial Reports used for ACF Non-Discretionary Grant Programs (0970–0510). The request revises ACF–696T, the annual financial report for tribal CCDF grant programs to submit financial claims. This specific form collects financial data for Tribal CCDF programs. The

proposed revisions to the ACF-696T will provide reporting instructions to Tribal Lead Agencies who are approved under a temporary opportunity to retroactively request the use CCDF funds, including most COVID-relief funds, for construction and/or major renovation with the intent of offsetting increased costs of materials, labor, and other related project costs.

DATES: *Comments due within 14 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above and below.

ADDRESSES: Copies of the proposed collection of information can be

obtained and comments may be forwarded by emailing infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: ACF programs require detailed financial information from their grantees that allows ACF to monitor various specialized cost categories within each program, to closely manage program activities, and to have sufficient financial information to enable periodic thorough and detailed audits. Generic Clearance for Financial Reports used for ACF Non-Discretionary Grant Programs allows ACF programs to efficiently develop and receive approval for financial reports that are tailored to

specific funding recipients and the associated needs of the program. For more information about the umbrella generic, see: https://www.reginfo.gov/public/do/PRAViewDocument?ref_nbr=202108-0970-002.

This specific GenIC collects financial data for tribal CCDF programs. The proposed revisions to the ACF-696T will provide reporting instructions to Tribal Lead Agencies who are approved under a temporary opportunity to retroactively request the use CCDF funds, including most COVID-relief funds, for construction and/or major renovation with the intent of offsetting increased costs of materials, labor, and other related project costs.

ANNUAL BURDEN ESTIMATES

Title of information collection	Number of respondents	Annual frequency of responses	Hourly burden per response	Annual hourly burden
ACF-696T	219	1	5	1,095

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden of the proposed collection of information; (c) 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. Consideration will be given to comments and suggestions submitted within 14 days of this publication.

Authority: 42 U.S.C. 9857, 42 U.S.C. 618.

Mary C. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2024-19916 Filed 9-4-24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2024-N-3902]

Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution and Related Collections of Information

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. 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 of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collections related to requirements for drug establishment registration and drug listing, including registrant reporting under the Federal Food, Drug, and Cosmetic Act (FD&C Act) with respect to listed drugs and certain guidances.

DATES: Either electronic or written comments on the collection of

information must be submitted by November 4, 2024.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of November 4, 2024. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.