document to ensure that the submitted CED study has a high degree of likelihood to generate the data and evidence needed to support Medicare coverage. *Form Number:* CMS–10697 (OMB control number 0938–1387); *Frequency:* Annually; *Affected Public:* Private sector—Not-for-profit institutions and Businesses or other for-profits; *Number of Respondents:* 13; *Total Annual Responses:* 13; *Total Annual Hours:* 1,300. (For policy questions regarding this collection, contact Andrew Ward at 4107–786–1794.)

**William N. Parham, III,**

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

[FR Doc. 2025–01242 Filed 1–16–25; 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; Objective Work Plan/On-going Progress Report (Office of Management and Budget #0970–0452)

**AGENCY:** Administration for Native Americans, Administration for Children and Families, United States Department of Health and Human Services.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration for Children and Families' (ACF) Administration for Native Americans (ANA) is requesting a 3-year extension to the Ongoing Progress Report (OPR) and the Objective Work Plan (OWP) (Office of Management and Budget (OMB) #0970–0452, expiration September 30, 2026). Changes are proposed only to the report.

**DATES:** *Comments due* February 18, 2025. OMB must decide 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 information in the OPR is collected on a semi-annual basis to monitor the performance of grantees and better gauge grantee progress.

The OPR information collection is conducted in accordance with Sec. 811 [42 U.S.C. 2992] of the Native American Programs Act and will allow ANA to report quantifiable results across all program areas. It also provides grantees with parameters for reporting their progress and helps ANA better monitor and determine the effectiveness of their projects.

The OWP information collection is conducted in accordance with 42 U.S.C. of the Native American Programs Act of 1972, as amended. This collection is necessary to evaluate applications for financial assistance and determine the relative merits of the projects for which such assistance is requested, as set forth in Section 806 [42 U.S.C. 2991–d 1](a)(1).

The report was revised based on a review by ANA and feedback from grantees, which identified some data elements that could be eliminated and areas that could be clarified.

*Respondents:* Federally and state recognized tribes, Native Pacific Islanders, Tribal Colleges and Universities, native non-profits, and consortia.

ANNUAL BURDEN ESTIMATES

| Instrument | Total number of respondents | Total number of responses per respondent | Average burden hours per response | Total burden hours | Annual burden hours |
|---|---|---|---|---|---|
| Objective Work Plan | 300 | 1 | 3 | 900 | 300 |
| Ongoing Progress Report | 200 | 6 | 1 | 1,200 | 400 |

*Estimated Total Annual Burden Hours:* 700.

*Authority:* Section 806 [42 U.S.C. 2991d–1](a)(1) and Section 811 [42 U.S.C. 2992].

**Mary C. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2025–01223 Filed 1–16–25; 8:45 am]

**BILLING CODE 4184–34–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Proposed Information Collection Activity; American Relief Act 2025 Disaster Supplemental Funds for Child Care (New Collection)

**AGENCY:** The 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 (OCC), Administration for Children and Families is requesting comment on the proposed application for disaster relief funding provided in the American Relief Act, 2025.

**DATES:** *Comments due* March 18, 2025. 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.

**ADDRESSES:** You can obtain copies of the proposed collection of information and submit comments by emailing *infocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The American Relief Act, 2025 provided $250,000,000 in disaster relief funding to OCC to distribute to

eligible States, territories, and Tribes in response to the consequences of major disasters and emergencies declared pursuant to the Robert T. Stafford Disaster Relief and Emergency Assistance Act (42 U.S.C. 5121 *et seq.*) occurring in 2023 and 2024.

OCC will be requesting information from eligible Child Care and Development Fund (CCDF) Lead Agencies who are interested in receiving these funds. The information requested includes the relevant major disaster or emergency declaration; a detailed description of the affected area; a detailed description of the impact on children, families, staff, and child care services; a description of each proposed activity; information on previous expenses incurred related to the disaster or emergency; and the total amount of funds requested. OCC will use the information received to inform decisions about distribution of funds.

*Respondents:* State, territory, and Tribal Lead Agencies.

**Annual Burden Estimates**

Respondents would provide one response to this request and information. The following burden estimates reflect the total estimated burden, which is expected within the first year of approval.

| Instrument | Total number of respondents | Total number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| ACF–OCC–CCDF–PI–2025–X (Disaster Supplemental Funds for Child Care—2023 and 2024 major disasters and emergencies) | 70 | 1 | 80 | 5,600 |

*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 60 days of this publication.

In addition, The Department solicits public comment on whether OCC should, in advance of receiving Lead Agency applications for funding, establish funding ranges for the size of grants based on certain factors, and if so, what those ranges and factors should be.

*Authority:* Public Law 118–158.

**Mary C. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2025–01221 Filed 1–16–25; 8:45 am]

**BILLING CODE 4184–87–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2025–N–0121]**

### Cell Therapies and Tissue-Based Products: A Public Workshop on Generating Scientific Evidence to Facilitate Development; Public Workshop; Request for Comments

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of public workshop; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA, the Agency, or we) is announcing the following virtual public workshop entitled ''Cell Therapies and Tissue-Based Products: A Public Workshop on Generating Scientific Evidence to Facilitate Development.'' The purpose of this workshop is to discuss the current state of the science for development and testing of certain cellular therapies and tissue-based products. In particular, FDA is convening this public workshop with relevant stakeholders to discuss best practices on generating scientific data necessary to further facilitate the development of cellular therapies, including stem cell products.

**DATES:** The public workshop will be held virtually on February 25, 2025, from 8:30 a.m. to 5 p.m. Eastern Time. Either electronic or written comments on this public workshop must be submitted by March 18, 2025. See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

**ADDRESSES:** The public workshop will be virtual.

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 March 18, 2025. 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.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

*Written/Paper Submissions*

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

*Instructions:* All submissions received must include the Docket No. FDA–2025–N–0121 for ''Cell Therapies and