

continue to use the ADR through the end of their grants.

The ADR 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.

*Respondents:* Tribal Government, Native non-profit organizations, and Tribal Colleges and Universities receiving ANA funding.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
ANA ADR .....	80	1	1	80

*Estimated Total Annual Burden Hours:* 80.

*Authority:* 42 U.S.C. 2992.

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for OMB Review; ACF-801: Child Care and Development Fund (CCDF) Quarterly Case-Level Report, (OMB #0970-0167)**

**AGENCY:** Office of Child Care, Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Office of Child Care (OCC), Administration for Children and Families (ACF) is requesting a 3-year extension of the form ACF-801: CCDF Quarterly Case-Level Report (OMB #0970-0167, expiration 2/28/2022). OCC proposes minor changes to the response categories under the following three data elements: Child’s gender, ethnicity, and race.

**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). All emailed requests should be identified by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The ACF-801 provides monthly case-level data on the children and families receiving direct child care services under CCDF. The ACF-801 case-level data are reported either monthly or quarterly. OCC added “no response” categories under the following three data elements: child’s gender, ethnicity, and race.

*Respondents:* State and Territory Lead Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
ACF-801: CCDF Quarterly Case—Level Report .....	56	4	25	5,600

*Estimated Total Annual Burden Hours:* 5,600.

*Authority:* Section 658K of the Child Care and Development Block Grant Act (42 U.S.C. 9858); regulations 45 CFR 98.70 and 98.71.

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**[Docket No. FDA-2013-N-1588]**

**Agency Information Collection Activities; Proposed Collection; Comment Request; Exemptions From Substantial Equivalence Requirements for Tobacco Products**

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) 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 exemptions from substantial equivalence requirements for tobacco products.