

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing infocollection@acf.hhs.gov. All 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.

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.

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.
 [FR Doc. 2021–26272 Filed 12–2–21; 8:45 am]
BILLING CODE 4184–81–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Numbers: 93.581, 93.587, 93.612]

Notice for Public Comment on Administration for Native Americans’ Program Policies and Procedures

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

ACTION: Request for public comment.

SUMMARY: Pursuant to Section 814 of the Native American Programs Act of 1974 (NAPA), as amended, the Administration for Native Americans (ANA) is required to provide members

of the public an opportunity to comment on proposed changes in interpretive rules and general statements of policy and to give notice of the proposed changes no less than 30 days before such changes become effective.

DATES: Comments are due by January 3, 2022. If ANA does not receive any significant comments within the 30-day comment period, ANA will proceed with the proposed changes in the respective published NOFOs. The NOFOs will serve as the final notice of these proposed changes.

ADDRESSES: Comments may be submitted to: Carmelia Strickland, Director of Program Operations, Administration for Native Americans, 330 C Street SW, Washington, DC 20201 or via email to: ANAComments@acf.hhs.gov.

FOR FURTHER INFORMATION CONTACT: Carmelia Strickland, Director, Division of Program Operations, Administration for Native Americans, 330 C Street SW, Washington, DC 20201; Telephone: (877) 922–9262; Email: ANAComments@acf.hhs.gov.

SUPPLEMENTARY INFORMATION: In accordance with notice requirements of NAPA, ANA herein describes proposed interpretive rules and general statements of policy that relate to ANA’s funding opportunities in Fiscal Year (FY) 2022. Changes to FY 2022 Notice of Funding Opportunity (NOFOs) will be based on the following previously published programs: Environmental Regulatory Enhancement (ERE), HHS–2021–ACF–ANA–NR–1907; Native American Language Preservation and Maintenance—Esther Martinez Immersion (EMI), HHS–2021–ACF–ANA–NB–1958; Native American Language Preservation and Maintenance (P&M), HHS–2021–ACF–ANA–NL–1924; Social and Economic Development Strategies (SEDS), HHS–2021–ACF–ANA–NA–1906; Social and Economic Development Strategies—

Alaska (SEDS–AK); and HHS–2021–ACF–ANA–NK–1902.

Section 814 of NAPA, as amended, (42 U.S.C. 2992b–1) incorporates provisions of the Administrative Procedure Act that require ANA to provide notice of its proposed interpretive rules and statements of policy and to seek public comment on such proposals. This notice serves to fulfill the statutory notice and public comment requirement. ANA voluntarily includes rules of practice and procedures in this notice in an effort to be transparent. The proposed interpretive rules, statements of policy, and rules of ANA practice and procedure reflected in clarifications, modifications, and new text will appear in the five FY 2022 NOFOs: ERE, EMI, P&M, SEDS, and SEDS–AK.

A. Interpretive rules, statements of policy, procedures, and practice. The proposals below reflect ANA’s proposed changes in rules, policy, or procedure that will take effect in the FY 2022 NOFOs.

1. Discontinuation of SEDS–GO

ANA has several new legislative economic development priorities under the Indian Community Economic Enhancement Act of 2020 (Public Law 116–261 Section 5), and Congress would like ANA to prioritize at least 50 percent of our available SEDS funding to go towards those types of projects. Therefore, ANA will discontinue SEDS–GO, and applicants can propose governance or organizational capacity building projects under regular SEDS.

2. Raising the Funding Level of SEDS–AK NOFO

Operating costs for grant-funded projects in Alaska are often higher than in the lower 48 states. In addition, ANA has seen a decline in the number of applications received over the last few years for the SEDS–AK funding competition. Therefore, in an effort create more interest in the program and to address the higher expenses, ANA

will increase the funding level for SEDS-AK from \$200,000 to \$300,000.

3. Clarification to EMI NOFO

In accordance with 42 U.S.C. 2991b-3(c)(7), applicants for an EMI grant must submit an official document that certifies the applicant has at least 3 years of experience in operating and administering a Native American language survival school, a Native American language nest, or any other educational program in which instruction is conducted in a Native American language. ANA has decided not to fund applicants that did not provide the certification as required by law. To reiterate and also clarify, the applicant must provide the required certification of having not less than 3 years of experience in operating and administering a Native American language survival school, a Native American language nest, or any other educational program in which instruction is conducted in a Native American language. The applicant may partner with other eligible entities (as defined under *Section III.1 Eligible Applicants* in the NOFO) that do not have to meet the certification requirement.

Statutory Authority: Section 814 of the Native American Programs Act of 1974 (NAPA), as amended.

Hope MacDonald LoneTree,

Deputy Commissioner, Administration for Native Americans.

[FR Doc. 2021-26271 Filed 12-2-21; 8:45 am]

BILLING CODE 4184-34-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0921]

Agency Information Collection Activities; Proposed Collection; Comment Request; Standards for the Growing, Harvesting, Packaging, and Holding of Produce for Human Consumption

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 the information collection associated with the standards for the growing, harvesting, packing, and holding of produce for human consumption.

DATES: Submit either electronic or written comments on the collection of information by February 1, 2022.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before February 1, 2022. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of February 1, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is 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-2011-N-0921 for "Agency Information Collection Activities; Proposed Collection; Comment Request; Standards for the Growing, Harvesting, Packaging, and Holding of Produce for Human Consumption." Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- **Confidential Submissions—**To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Dockets Management