ANNUAL BURDEN ESTIMATES—Continued

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Interview guide for PHA administrator and FUP liaison	16	8	1	1	8
Focus group guide for youth	96	48	1	1.5	72
Administrative data list	24	12	1	5	60
Total					624

Estimated Total Annual Burden Hours: 624.

Authority: Title IV–E of the Social Security Act, IV–E § 477(g)(1–2), as amended by the Foster Care Independence Act of 1999.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2020–02315 Filed 2–5–20; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.587]

Notice for Public Comment on Administration for Native Americans' Program Policies Relating to the Native American Language Preservation and Maintenance—Esther Martinez Immersion Funding Opportunity

AGENCY: Administration for Native Americans (ANA), Administration for Children and Families (ACF), Department of Health and Human Services (HHS).

ACTION: Notice for public comment.

SUMMARY: Pursuant to Section 814 of the Native American Programs Act of 1974 (NAPA), as amended, the 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. In accordance with the notice requirements of NAPA, ANA herein describes proposed general statements of policy that relate to the Native American Language Preservation and Maintenance—Esther Martinez Immersion (EMI) funding opportunity announcement (FOA) in fiscal year (FY) 2020.

DATES: Comments are due by March 9, 2020. If ANA does not receive any significant comments on the statements of policy within the 30-day comment period, ANA will proceed with the proposed changes in the published FOA. The FOA 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: *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: 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. ANA voluntarily provides notices of its rules of practice and procedure in an effort to be transparent. ANA published a notice in the **Federal Register** on December 27, 2019, located at 84 FR 71428, pages 71428–71430, that lays out changes ANA intends to implement in its FOAs for FY 2020. This notice provides information on additional changes that will be implemented in the FY 2020 EMI FOA.

ANA's past FOAs can be accessed at *http://www.acf.hhs.gov/grants/open/foa/office/ana* or *http://www.acf.hhs.gov/grants/open/foa/.* Synopses and application forms will be available on *https://www.grants.gov.*

1. On December 20, 2019, President Trump signed into law the Esther Martinez Native American Languages Programs Reauthorization Act (Reauthorization Act). The statute amended NAPA to allow ANA to set the project period of EMI grants to up to 60 months. Because of the complexity of language grants that provide immersion instruction and because Native American communities are most knowledgeable about their language programs and the length of time necessary to implement them, ANA has determined to use this new authority to give tribes maximum flexibility on the length of their grant and allow the project period for the FY 2020 EMI grants to be up to 60 months. Previously, awards were limited to a 36-month project period. FY 2020 awards will be made for a project period of 36 months, 48 months, or 60 months. An initial award will be for a 12-month budget period. Non-competing continuation (NCC) awards will be awarded on the basis of annual NCC applications, availability of funds, satisfactory progress, on-time completion of grant reporting requirements, and a determination that continued funding would be in the best interest of the government.

2. ANA had intended to accept applications for the EMI grants for 90 days, in-line with its other grants. However, because ANA is making additional changes to the EMI grant to implement the Reauthorization Act, ANA will reduce the period for accepting applications in order to issue the EMI awards in a timely manner. The EMI FOA, HHS–2018–ACF–ANA–NB–1343, will accept applications for 60 days from the date it is published on *Grants.gov*.

3. ANA plans to change the order of the publication of the FOAs. In the Notice of Public Comment published on December 27, 2019, 84 FR 71428, pages 71428–71430, Supplementary Information, ANA stated that it planned to publish the FOAs for the EMI, Native American Language Preservation and Maintenance (P&M), and Environmental Regulatory Enhancement (ERE) grants first and then allow a 2-week period before the Social and Economic Development Strategies (SEDS) and Social and Economic Development Strategies-Alaska (SEDS–AK) FOAs were published. However, because of the changes to the EMI grant described above, ANA will need additional time to publish the EMI FOA. In order to continue to stagger the submission deadlines, ANA has decided it will publish the P&M, ERE, and SEDS–AK FOAs first and then allow a 2-week period before the SEDS and EMI FOAs are published. As this is a change in

agency procedure, the change will be made without reviewing comments.

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

Elizabeth Leo,

Senior Grants Policy Specialist, Division of Grants Policy, Office of Administration.

[FR Doc. 2020–02319 Filed 2–5–20; 8:45 am]

BILLING CODE 4184–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0118]

Agency Information Collection Activities; Proposed Collection; Comment Request; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002

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 collection provisions of our regulations requiring that the Agency receive prior notice before food is imported or offered for import into the United States.

DATES: Submit either electronic or written comments on the collection of information by April 6, 2020.

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 April 6, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of April 6, 2020. 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–2010–N–0118 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002.'' 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.

- 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 Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3521), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''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 (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an