

*Response:* Thank you for your comment. The new ICEEA law does allow for the development of existing Native CDFIs. Therefore, should a Native CDFI submit an application that proposes a project for any of the following projects: (1) The development of a tribal code or courts system for purposes of economic development, including commercial codes, training for court personnel, (2) the development of non-profit subsidiaries or other tribal business structures; or “(3) the development of a tribal master plan for community and economic development and infrastructure” and the application includes the economic priority area(s) in the project goal, all objectives and indicators as reflected in the project’s framework, project approach, OWP, and outcome tracker, they will be awarded points. ANA will instruct reviewers to provide all bonus points for applications that propose an economic priority project that expands or creates a Native CDFI.

**Elizabeth Leo,**

*Senior Grants Policy Specialist, Office of Grants Policy, Administration for Children and Families.*

[FR Doc. 2021-16959 Filed 8-9-21; 8:45 am]

**BILLING CODE 4184-34-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

#### Submission for OMB Review; Survey of the National Survey of Child and Adolescent Well-Being (NSCAW) Adopted Youth, Young Adults, and Adoptive Parents (0970-0555)

**AGENCY:** Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration for Children and Families (ACF) at the U.S. Department of Health and Human Services (HHS) seeks approval for an extension with no changes to a one-time study to examine familial outcomes 8 years or more after a child’s adoption from the child welfare system. The primary objective of this study is to estimate the prevalence of instability events that occur in families who have adopted children who have exited the foster care system. The second objective is to understand risk and protective factors associated with post adoption instability. Office of Management and Budget (OMB) approval expires September 30, 2021, and this request is to extend approval to allow for the completion of data collection.

**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.

#### SUPPLEMENTARY INFORMATION:

**Description:** Through this study, ACF is conducting web or telephone surveys with adopted youth, young adults, and adults as well as adoptive parents who were participants in the first or second cohort of NSCAW (NSCAW I, II; OMB #0970-0202). The surveys are designed to collect information about instability events (such as foster care re-entry or running away that occurred after a child’s adoption) as well as family functioning, perceptions of the adoption relationship, and services and support received after adoption. Due to the COVID-19 pandemic, initial activities to contact potential respondents were delayed. As a result, ACF is requesting an extension to collect data beyond the current OMB expiration date of September 30, 2021.

**Respondents:** Adopted youth, young adults, adults, and their associated adoptive parents who participated in NSCAW I or II.

### ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Survey of NSCAW Adopted Youth, Young Adults, and Adults .....	588	1	.5	294	294
Survey of NSCAW Adoptive Parents .....	554	1	.5	277	277

*Estimated Total Annual Burden Hours:* 571.

**Authority:** Child Abuse Prevention and Treatment and Adoption Reform Act of 1978.

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2021-16979 Filed 8-9-21; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket Nos. FDA-2010-N-0190; FDA-2012-N-0197; FDA-2014-N-1414; and FDA-2014-N-0913]

#### Agency Information Collection Activities; Announcement of Office of Management and Budget Approvals

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is publishing a list of information collections that have been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD

20852, 301–796–7726, [PRAStaff@fda.hhs.gov](mailto:PRAStaff@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** The following is a list of FDA information collections recently approved by OMB under section 3507 of the Paperwork

Reduction Act of 1995 (44 U.S.C. 3507). The OMB control number and expiration date of OMB approval for each information collection are shown in table 1. Copies of the supporting statements for the information collections are available on the internet

at <https://www.reginfo.gov/public/do/PRAMain>. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

TABLE 1—LIST OF INFORMATION COLLECTIONS APPROVED BY OMB

Title of collection	OMB control number	Date approval expires
Infant Formula Requirements .....	0910–0256	5/31/2024
Shortages Data Collection .....	0910–0491	6/30/2024
Guidance on Labeling for Natural Rubber Latex Condoms .....	0910–0633	6/30/2024
Section 513(g) Requests for Information .....	0910–0705	6/30/2024

Dated: August 5, 2021.

**Lauren K. Roth,**

*Acting Principal Associate Commissioner for Policy.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA–2020–N–1648]

#### **Pediatric Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments**

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

**ACTION:** Notice; establishment of a public docket; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Pediatric Advisory Committee. The general function of the committee is to provide advice and recommendations to FDA on regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comment on this document.

**DATES:** The meeting will be held on September 17, 2021, from 10 a.m. to 4:30 p.m. Eastern Time.

**ADDRESSES:** Please note that due to the impact of the COVID–19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: <https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm>.

FDA is establishing a docket for public comment on this meeting. The docket number is FDA–2020–N–1648. The docket will close on September 16, 2021. Submit either electronic or written comments on this public meeting by September 16, 2021. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before September 16, 2021. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of September 16, 2021. 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.

Comments received on or before September 3, 2021, will be provided to the committee. Comments received after that date will be taken into consideration by FDA. In the event that the meeting is cancelled, FDA will continue to evaluate any relevant applications or information, and consider any comments submitted to the docket, as appropriate.

You may submit comments as follows:

#### *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–2020–N–1648 for “Pediatric Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments.” 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