## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review: ACF Program Instruction—Children's Justice Act (OMB #0970–0425)

**AGENCY:** Children's Bureau; Administration for Children and Families; HHS

**ACTION:** Request for Public Comment.

**SUMMARY:** The Administration for Children and Families (ACF) is requesting a 3-year extension of the Children's Justice Act Program Instruction (OMB #0970–0425, expiration 4/30/2020). There are no changes requested to the form.

**DATES:** *Comments due within 30 days of publication.* OMB is required to make a decision concerning 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.

**SUPPLEMENTARY INFORMATION:**

*Description:* The Program Instruction, prepared in response to the enactment of the Children's Justice Act (CJA), Title II of Public Law 111–320, Child Abuse Prevention and Treatment Act Reauthorization of 2010, provides direction to the states and territories to accomplish the purposes of assisting states in developing, establishing, and operating programs designed to improve: (1) The assessment and investigation of suspected child abuse and neglect cases, including cases of suspected child sexual abuse and exploitation, in a manner that limits additional trauma to the child and the child's family; (2) the assessment and investigation of cases of suspected child abuse-related fatalities and suspected child neglect-related fatalities; (3) the investigation and prosecution of cases of child abuse and neglect, including child sexual abuse and exploitation; and (4) the assessment and investigation of cases involving children with disabilities or serious health-related problems who are suspected victims of child abuse or neglect. This Program Instruction contains information collection requirements that are found in Public Law 111–320 at sections 107(b) and 107(d), and pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute; to monitor, evaluate, and measure grantee achievements in addressing the investigation and prosecution of child abuse and neglect; and to report to Congress.

*Respondents:* State Governments.

ANNUAL BURDEN ESTIMATES

| Instrument | Total number of respondents | Annual number of responses per respondent | Average burden hours per response | Annual burden hours |
|---|---|---|---|---|
| Children's Justice Act Program Instruction | 52 | 1 | 60 | 3,120 |

*Estimated Total Annual Burden Hours:* 3,120.

**Authority:** 42 U.S.C. 5106c Sec. 107 (b)(4) and 42 U.S.C. 5106 Sec. 107 (B)(5).

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020–06525 Filed 3–27–20; 8:45 am]

**BILLING CODE 4184–25–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Proposed Information Collection Activity; Survey of the National Survey of Child and Adolescent Well-Being (NSCAW) Adopted Youth, Young Adults, and Adoptive Parents (New Collection)

**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 a one-time study to examine familial outcomes 8 or more years 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.

**DATES:** *Comments due within 60 days of publication.* In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

**ADDRESSES:** Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing *OPREinfocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests emailed or written should be identified by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The proposed study would conduct 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.

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

ANNUAL BURDEN ESTIMATES

| Instrument | Total/annual number of respondents | Number of responses per respondent | Average burden hours per response | Annual burden hours |
|---|---|---|---|---|
| Survey of NSCAW Adopted Youth, Young Adults, and Adults | 588 | 1 | .5 | 294 |
| Survey of NSCAW Adoptive Parents | 554 | 1 | .5 | 277 |

*Estimated Total Annual Burden Hours:* 571

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

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020–06491 Filed 3–27–20; 8:45 am]

**BILLING CODE 4184–25–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2020–N–0008]**

### Allergenic Products Advisory Committee; Cancellation of Meeting

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

**ACTION:** Notice.

**SUMMARY:** The meeting of the Allergenic Products Advisory Committee scheduled for May 15, 2020, is canceled. The Allergenic Products Advisory Committee meeting scheduled for May 15, 2020, to discuss and make recommendations on the safety and efficacy of Peanut (*Arachis hypogaea*) Allergen Extract manufactured by DBV Technologies, S.A, has been canceled to allow time for the FDA to review outstanding issues. The Agency intends to continue evaluating the product and will schedule an Advisory Committee meeting in the future, as needed. The meeting was announced in the **Federal Register** on February 24, 2020.

**FOR FURTHER INFORMATION CONTACT:** Kathleen Hayes, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6307, Silver Spring, MD 20993–0002, 301–796–7864, *kathleen.hayes@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting, which was announced in the **Federal Register** of February 24, 2020, 85 FR 10451.

Dated: March 24, 2020.

**Lowell J. Schiller,**

*Principal Associate Commissioner for Policy.*

[FR Doc. 2020–06513 Filed 3–27–20; 8:45 am]

**BILLING CODE 4164–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2016–N–0736]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Tracking Network for PETNet, LivestockNet, and SampleNet

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Fax written comments on the collection of information by April 29, 2020.

**ADDRESSES:** To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, Fax: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0680. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, *PRAStaff@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

### Tracking Network for PETNet, LivestockNet, and SampleNet

### OMB Control Number 0910–0680—Extension

The Center for Veterinary Medicine and the Partnership for Food Protection developed a web-based tracking network (the tracking network) to allow Federal, State, and Territorial regulatory and public health Agencies to share safety information about animal food. Information is submitted to the tracking network by regulatory and public health Agency employees with membership rights. The efficient exchange of safety information is necessary because it improves early identification and evaluation of a risk associated with an animal food product. We use the information to assist regulatory Agencies to quickly identify and evaluate a risk and take whatever action is necessary to mitigate or eliminate exposure to the risk. Earlier identification and communication with respect to emerging safety information may also mitigate the potential adverse economic impact for the impacted parties associated with such safety issues. The tracking network was developed under the requirements set forth under section 1002(b) of the Food and Drug Administration Amendments Act of 2007 (FDAAA) (Pub. L. 110–085). Section 1002(b) of the FDAAA required FDA, in relevant part, to establish a pet food early warning alert system.