emailed or written, should be identified by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* Since 2006, CB has awarded multiple rounds of competitive grants to state and local agencies and service providers under the RPG Program. Grants are awarded to organizations such as child welfare agencies, substance abuse treatment providers, or family court systems to develop interagency collaborations and provide services designed to increase well-being, improve permanency, and enhance the safety of children who are in or are at risk of being placed in out-of-home care as a result of a parent's or caretaker's substance abuse. Thirty-five grantees are participating in the ongoing RPG national cross-site evaluation, which examines implementation, partnerships, outcomes, and impacts. All grantees collect data on a uniform set of performance measures and report them to CB on a semi-annual basis through a web-based system. These ongoing data collection activities are approved under OMB #0970–0527. All grantees are also required to use a portion of their funding to conduct their own ''local'' program impact evaluation.

This proposed cost study adds a new and unique contribution to CB's portfolio of evaluation activities. Although the RPG cross-site evaluation will provide evidence for the effectiveness of some interventions to address the emotional effects of trauma, more information is needed about the cost of implementing these Evidence-Based Programs (EBPs).

The cost study has the key objective to determine the cost of implementing three select Trauma-Specific EBPs: Parent-Child Interaction Therapy, Seeking Safety, and Trauma-Focused Cognitive Behavioral Therapy. To carry out this objective, the study team will collect detailed cost information from nine RPG round four and five grantees who are implementing these selected EBPs. For each grantee, the study team will administer two data collection instruments: (1) A Cost Workbook used to collect comprehensive information on the cost of implementing each select program (Instrument #1), and (2) a Staff Survey and Time Log used to collect information on how program staff allocate their time (Instrument #2).

*Respondents:* Grantee staff.

**Annual Burden Estimates**

Data collection will take place within a 1-year period.

| Instrument | Total number of respondents | Total number of responses per respondent | Average burden hours per response | Total/annual burden hours |
|---|---|---|---|---|
| Cost Workbook | 9 | 1 | 8 | 72 |
| Staff Survey and Time Log | 90 | 1 | 3.6 | 330 |

*Estimated Total Annual Burden Hours:* 402.

*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:** The Child and Family Services Improvement and Innovation Act (Pub. L. 112–34).

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020–13935 Filed 6–26–20; 8:45 am]

**BILLING CODE 4184–29–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 (New Collection)

**AGENCY:** Office of Planning, Research and Evaluation (OPRE); Administration for Children and Families; Department Of Health and Human Services (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 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 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 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.

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

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020–13939 Filed 6–26–20; 8:45 am]

**BILLING CODE 4184–25–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Community Living

### Notice of Intent To Award a Sole Source Supplement to the Christopher and Dana Reeve Foundation

**ACTION:** Notice of Intent to award a sole source supplement.

**SUMMARY:** The Administration for Community Living (ACL) is announcing the award of single-source supplement for the National Paralysis Resource Center (PRC) that was included in the 2020 Congressional budget appropriations.

**SUPPLEMENTARY INFORMATION:** The National Paralysis Resource Center is operated by the Christopher and Dana Reeve Foundation, which offers important programmatic opportunities for persons with disabilities and older adults. The PRC provides comprehensive information for people living with spinal cord injury, paralysis, and mobility-related disabilities and their families. Resources include information and referral by phone and email in multiple languages; a peer and family support mentoring program; a military and veterans program; multicultural outreach services; multiple quality of life grants; and a national website. The administrative supplement for FY2020 will be in the amount of $2,188,339, bring the total award for FY20 to $8,700,000.

*Program Name:* National Paralysis Resource Center.

*Recipient:* Christopher and Dana Reeve Foundation.

*Period of Performance:* The supplement award will be issued for the third year of a three year project a project period, July 1, 2020 through June 30, 2021.

*Award Amount:* $2,188,339.

*Award Type:* Cooperative Agreement.

*Statutory Authority:* This program is authorized under Section 317 of the Public Health Service Act (42 U.S.C. 247(b–4)).

*CFDA Number:* 93.325 Discretionary Projects.

The purpose of the supplemental funding is to support the expansion the National Paralysis Resource Center to improve the health and quality of life of individuals living with paralysis and their families by raising awareness of and facilitating access to a broad range of services relevant to individuals with paralysis. With the additional funding, the PRC will work to expand the National Resource and Information Center; increase the health and quality of life of Americans with disabilities living with paralysis; increase support and resources to people with paralysis, their families and caregivers; expand collaboration with federal agencies and other national organizations that have a vested interest in the paralysis community; and strengthen performance measures.

Dated: June 17, 2020.

**Mary Lazare,**

*Principal Deputy Administrator, Administration for Community Living.*

[FR Doc. 2020–13577 Filed 6–26–20; 8:45 am]

**BILLING CODE 4154–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2019–D–0914]**

### Review and Update of Device Establishment Inspection Processes and Standards; Guidance for Industry; Availability

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

**ACTION:** Notice of availability.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance entitled ''Review and Update of Device Establishment Inspection Processes and Standards.'' FDA is issuing this guidance to comply with changes to the Federal Food, Drug, and Cosmetic Act (FD&C Act) as amended by the FDA Reauthorization Act of 2017 (FDARA), which requires that FDA review and update, as needed, the processes and standards applicable to inspections (other than for-cause) of domestic and foreign medical device establishments in place as of August 18, 2017. This guidance describes how FDA will implement uniform inspection processes and standards. The guidance also describes standardized methods of communication during the inspection process and identifies practices for investigators and device establishments to facilitate the continuity of inspections of such establishments.

**DATES:** The announcement of the guidance is published in the **Federal Register** on June 29, 2020.

**ADDRESSES:** You may submit either electronic or written comments on Agency guidances at any time 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