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