of the designated State entity, and not less than 51 percent of the directors of the centers for independent living in the State. ACL reviews the SPIL for compliance with the Rehabilitation Act and 45 CFR part 1329 and approves it. The SPIL also serves as a primary planning document for continuous monitoring of, and technical assistance to, the state independent living programs to ensure appropriate planning, financial support and coordination, and other assistance to appropriately address, on a statewide basis, needs for the provision of independent living services in the state.

The proposed data collection tools may be found on the ACL website for review at *https://www.acl.gov/about-acl/public-input*.

Estimated Program Burden: ACL estimates the burden of this collection of information as follows: 56 Statewide Independent Living Councils will respond to the requirement for a SPIL every three years. It will take approximately 60 hours for each state's Statewide Independent Living Council to jointly complete the development of the SPIL for a total of approximately 3,360 hours. This estimate is based on amounts of time that Statewide Independent Living Councils have reported that they have spent responding to previous requests for this report. ACL is not requesting any change in the data States are required to submit. As such, there is no change to the estimated reporting burden.

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
Statewide Independent Living Councils	56	1	60	3,360
Total	56	1	60	3,360

Dated: October 10, 2018.

Mary Lazare,

Principal Deputy Administrator.

[FR Doc. 2018–22753 Filed 10–18–18; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Public Comment Request; Centers for Independent Living Program Performance Report (0985–NEW)

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to comment on the proposed collection of information listed above. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish a 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 New Data Collection (ICR New) solicits comments on the information collection requirements relating to the Centers for Independent Living *under* the Rehabilitation Act of 1973.

DATES: Comments on the collection of information must be submitted electronically by 11:59 p.m. (EST) or postmarked by December 18, 2018.

ADDRESSES: Submit electronic comments on the collection of information to: Peter Nye at *peter.nye@acl.hhs.gov*. Submit written comments on the collection of information to Administration for Community Living, Washington, DC 20201, Attention: Peter Nye.

FOR FURTHER INFORMATION CONTACT: Peter Nye, Administration for Community Living, Washington, DC 20024, (202) 795–7606 or *peter.nye@acl.hhs.gov*.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), 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 existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing a notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, ACL invites comments on our burden estimates or any other aspect of this collection of information, including:

(1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility;

(2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used to determine burden estimates;

(3) ways to enhance the quality, utility, and clarity of the information to be collected; and

(4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

In the context of ACL, IL programs are supported through funding authorized by the Rehabilitation Act of 1973, as amended (The Act). Title VII, chapter 1 of the Act states the current purpose of the program is to ''promote a philosophy of independent living including a philosophy of consumer control, peer support, self-help, self-determination, equal access, and individual and system advocacy, in order to maximize the leadership, empowerment, independence, and productivity of individuals with disabilities, and the integration and full inclusion of individuals with disabilities into the mainstream of American society.''

ILS PPR and CIL PPR are being submitted separately because they are separate collections of different information from different parties. Separating these PRA processes reduces confusion and increases the Independent Living Administration's ability to identify issues specific to CILs. This request is for CIL PPR, which is submitted annually by all CILs receiving IL Subchapter C funds. The PPRs are used by ACL to assess grantees' compliance with title VII of the Act, and with 45 CFR 1329 of the Code of Federal

Regulations and with applicable provisions of the HHS Regulations at 45 CFR part 75. The PPR serves as the primary basis for ACL's monitoring activities in fulfillment of its responsibilities under sections 706 and 722 of the Act. The PPR also enables ACL to track performance outcomes and efficiency measures of the Centers for Independent Living (CIL) programs with respect to the annual and long-term performance targets established in compliance with GPRA. The PPR is also used by ACL to design CIL and SILC training and technical assistance programs authorized by section 711A and section 721 of the Act.

ACL published a **Federal Register** Notice regarding the independent living programs information collection on February 23, 2017. Two-hundred and twenty-one individual comments were received. The responses indicated a need to make substantial changes to the collection. The current version of the ILS PPR that ILA is requesting an extension for was approved by OMB, but will expire on December 31, 2018. Further deliberation is needed to ensure that we appropriately address all of the concerns. ILA is proposing to extend the currently approved forms for three years while we work on a revision that addresses all the suggested changes.

The proposed data collection tools may be found on the ACL website for review at *https://www.acl.gov/about-acl/public-input*.

Estimated Program Burden: ACL estimates the burden of this collection of information as follows: 353 Centers for Independent Living will each complete one CIL PPR annually, and it will take an estimated 35 hours per CIL for an estimated total of 12,355 hours. This burden estimate is based partly on ILA's estimates of how long CILs probably take to find the information that PPRs ask for and partly on what CILs have told the Independent Living Administration about how long filling out the PPRs took.

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
Centers for Independent Living	353	1	35	12,355

Dated: October 9, 2018.

Mary Lazare,

Principal Deputy Administrator.

[FR Doc. 2018–22754 Filed 10–18–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–1072]

Agency Information Collection Activities; Proposed Collection; Comment Request; Application for Participation in Food and Drug Administration Fellowship and Traineeship Programs

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) 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 ''Application for Participation in FDA Fellowship and Traineeship Programs.''

DATES: Submit either electronic or written comments on the collection of information by December 18, 2018.

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 December 18, 2018. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of December 18, 2018. 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–2014–N–1072 for ''Application for Participation in Food and Drug Administration Fellowship and Traineeship Programs.'' 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