

license), address, organization, telephone number(s), fax number, and email address. You will receive a registration confirmation with instructions for your arrival at the CMS complex or you will be notified that the seating capacity has been reached.

IV. Security, Building, and Parking Guidelines

This meeting will be held in a federal government building; therefore, federal security measures are applicable. The Real ID Act, enacted in 2005, establishes minimum standards for the issuance of state-issued driver's licenses and identification (ID) cards. It prohibits Federal agencies from accepting an official driver's license or ID card from a state unless the Department of Homeland Security determines that the state meets these standards. Beginning October 2015, photo IDs (such as a valid driver's license) issued by a state or territory not in compliance with the Real ID Act will not be accepted as identification to enter Federal buildings. Visitors from these states/territories will need to provide alternative proof of identification (such as a valid passport) to gain entrance into CMS buildings. The current list of states from which a Federal agency may accept driver's licenses for an official purpose is found at <http://www.dhs.gov/real-id-enforcement-brief>. We recommend that confirmed registrants arrive reasonably early, but no earlier than 45 minutes prior to the start of the meeting, to allow additional time to clear security. Security measures include the following:

- Presentation of government-issued photographic identification to the Federal Protective Service or Guard Service personnel.
- Inspection of vehicle's interior and exterior (this includes engine and trunk inspection) at the entrance to the grounds. Parking permits and instructions will be issued after the vehicle inspection.
- Inspection, via metal detector or other applicable means, of all persons entering the building. We note that all items brought into CMS, whether personal or for the purpose of presentation or to support a presentation, are subject to inspection. We cannot assume responsibility for coordinating the receipt, transfer, transport, storage, set-up, safety, or timely arrival of any personal belongings or items used for presentation or to support a presentation.

Note: Individuals who are not registered in advance will not be permitted to enter the building and will be unable to attend the

meeting. The public may not enter the building earlier than 45 minutes prior to the convening of the meeting.

All visitors must be escorted in areas other than the lower and first floor levels in the Central Building.

V. Collection of Information

This document does not impose information collection requirements, that is, reporting, recordkeeping or third-party disclosure requirements. Consequently, there is no need for review by the Office of Management and Budget under the authority of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

Authority: 5 U.S.C. App. 2, section 10(a).

Dated: June 4, 2018.

Kate Goodrich,

Director, Center for Clinical Standards and Quality, Chief Medical Officer, Centers for Medicare & Medicaid Services.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Public Comment Request; State Councils on Developmental Disabilities—Annual Program Performance Report (PPR) (OMB Control Number—0985-0033)

AGENCY: Administration on Intellectual and Developmental Disabilities (AIDD), Administration for Community Living (ACL), U.S. Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-day notice collects comments on the information collection requirements related to the State Councils on Developmental Disabilities—Annual Program Performance Report (PPR) [Proposed Extension with Changes of a Currently Approved Collection (ICR Rev)].

DATES: Submit written comments on the collection of information by July 16, 2018.

ADDRESSES: Submit written comments on the collection of information by:

(a) Email to: OIRA_submission@omb.eop.gov, Attn: OMB Desk Officer for ACL;

(b) fax to 202.395.5806, Attn: OMB Desk Officer for ACL; or

(c) by mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

FOR FURTHER INFORMATION CONTACT: Sara Newell-Perez at (202) 795-7413 or Sara.Newell-Perez@acl.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, ACL has submitted the following proposed collection of information to OMB for review and clearance. The proposed data collection represents a revision of a currently approved information collection (ICR-Rev). This collection is necessary for the proper performance and function of the agency. On an annual basis, Councils are required to submit a Program Performance Report (PPR) to describe the extent to which annual progress is being achieved on the 5-year State plan goals. The PPR will be used by (1) the Council as a planning document to track progress made in meeting state plan goals; (2) the citizenry of the State as a mechanism for monitoring progress and activities on the plans of the Council; and (3) the Department as a stewardship tool for ensuring compliance with the Developmental Disabilities Assistance and Bill of Rights Act of 2000 and for monitoring and providing technical assistance (e.g., during site visits), and support for management decision making.

Comments in Response to the 60-Day Federal Register Notice

A notice was published in the **Federal Register** on October 4, 2017 (Vol. 82, Number 191; pp. 46246-46247). Two comments were received. The first was a comment about ACL and policies around deinstitutionalization. The second comment requested that Councils have more transparency and make their PPRs available to the public via their websites. ACL appreciates and understands these comments. Although ACL recognizes that these comments might provide useful information for the program, it is not required to meet the statutory requirements for this program. No change is proposed.

The proposed template may be found on the ACL website at <https://www.acl.gov/about-acl/public-input>.

Estimated Program Burden

ACL estimates the burden of this collection of information as follows: The

total estimated hour burden per respondent for the proposed DD Council PPR will increase from the 138 hours

estimated in 2015 to 172 burden hours per response. The number of hours is multiplied by 56 State Council

programs, resulting in a total estimated hour aggregate burden of 9,632.

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
State Councils on Developmental Disabilities/Program Performance Report	56	1	172	9632
Total	56	1	172	9632

Dated: June 7, 2018.
Lance Robertson,
Administrator and Assistant Secretary for Aging.
 [FR Doc. 2018-12826 Filed 6-14-18; 8:45 am]
BILLING CODE 4154-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Notice of Intent To Award a Single-Source Supplement

ACTION: Intent To Award a Single-Source Supplement for the Amputee Coalition of America, Inc. for the National Limb Loss Resource Center Cooperative Agreement.

The Administration for Community Living (ACL) announces the intent to award a single-source supplement to the current cooperative agreement held by the Amputee Coalition of America, Inc. for the National Limb Loss Resource Center (NLLRC). The purpose of this project is to expand on current grant activities, such as increasing activities and programs that promote health, wellness, and the adoption of healthy behaviors with the objective of preventing and/or reducing chronic conditions associated with limb loss and increase partnerships and collaborations with ACL programs that will benefit all people living with limb loss or limb differences. The administrative supplement for FY 2018 will be in the amount of \$669,905, bringing the total award for FY 2018 to \$3,697,142.

The additional funding will not be used to begin new projects. The funding will be used to enhance and expand existing programs that can serve an increased number of veterans and people living with limb loss and limb differences by providing increased technical assistance activities; promoting health and wellness programs; promoting the adoption of healthy behaviors with the objective of preventing and/or reducing chronic

conditions associated with limb loss; increasing partnerships and collaborations with ACL programs that will benefit all people living with limb loss or limb differences; enhancing and expanding the evaluation activities currently under way; and enhancing website capacities for improved information dissemination.

Program Name: National Limb Loss Resource Center.

Recipient: The Amputee Coalition of America, Inc.

Period of Performance: The supplement award will be issued for the third year of the three-year project period of April 1, 2016, through March 29, 2019.

Total Award Amount: \$669,905 in FY 2018.

Award Type: Cooperative Agreement Supplement.

Statutory Authority: This program is authorized under Section 317 of the Public Health Service Act (42 U.S.C. 247(b-4)); Consolidated and Further Continuing Appropriations Act, 2015, Public Law 113-235 (Dec. 16, 2014).

Basis for Award: The Amputee Coalition of America, Inc. is currently funded to carry out the objectives of this program, entitled *The National Limb Loss Resource Center* for the period of April 1, 2016, through March 29, 2019. Since the program transferred from CDC to ACL in late 2015, the grantee has accomplished a great deal. The supplement will enable the grantee to carry their work even further, serving more people living with limb loss and/or limb differences and providing even more comprehensive training and technical assistance in the development of LTSS supportive services. The additional funding will not be used to begin new projects or activities. The NLLRC will enhance and expand currently funded activities such as conducting national outreach for the development and dissemination of patient education materials, programs, and services; providing technical support and assistance to community based limb loss support groups; and raising awareness about the limb loss and limb differences communities.

Establishing an entirely new grant project at this time would be potentially disruptive to the current work already well under way. More importantly, the people living with limb loss and limb differences currently being served by this program could be negatively impacted by a service disruption, thus posing the risk of not being able to find the right resources that could negatively impact on health and wellbeing. If this supplement were not provided, the project would be less able to address the significant unmet needs of additional limb loss survivors. Similarly, the project would be unable to expand its current technical assistance and training efforts in NLLRC concepts and approaches, let alone reach beyond traditional providers of services to this population to train more “mainstream” providers of disability services.

For More Information Contact: For further information or comments regarding this program supplement, contact Elizabeth Leef, U.S. Department of Health and Human Services, Administration for Community Living, Administration on Disabilities, Independent Living Administration: telephone (202) 475-2486; email Elizabeth.leef@acl.hhs.gov.

Dated: June 6, 2018.
Lance Robertson,
Administrator and Assistant Secretary for Aging.
 [FR Doc. 2018-12978 Filed 6-14-18; 8:45 am]
BILLING CODE 4154-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2018-D-1788]

Intravascular Catheters, Wires, and Delivery Systems With Lubricious Coatings—Labeling Considerations; Draft Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.