equipment, etc.) of increasing end-user compliance with proper selection, care, maintenance, and use of PPT; (10) provides systematic collection, analysis, and interpretation of PPT use practices, including investigation of barriers to effective PPT use; (11) produces and disseminates technical information, research findings, training materials, and recommendations for PPT to improve protection of workers; (12) evaluates and disseminates PPT performance trends published through the post market surveillance activities; and (13) identifies and implements an effective communication and outreach program for stakeholders within the NIOSH sectors to inform end users of proper selection, care, maintenance, and use of PPT.

Conformity Verification and Standards Development Branch (CCLG). (1) Administers the Department of Health and Human Services Title 42 Code of Federal Regulations (CFR), Part 84-Respiratory Protective Devices conformity assessment functions (*i.e.* inspection, testing, certification, documentation control, quality assurance, and surveillance) including: (a) Processing respirator approval applications by verifying conformance with Federal regulations and national consensus standards such as performance, quality, reliability, and documentation requirements to determine the effectiveness of respirators used during entry into or escape from hazardous atmospheres, (b) issuing or revoking NIOSH certificates of approval, (c) evaluating and maintaining official records on NIOSH-certified respirators including the establishment of NPPTL and national databases, (d) recommending NIOSH policy relating to RPD conformity verification criteria for traditional and innovative respirator technologies and applications, and, (e) investigating and processing Freedom-of-Information-Act requests; (2) establishes and administers an internal audit program to evaluate the conformity assessment functions of NPPTL; (3) maintains official files of policies, standards, standard operating and test procedures used as the basis for granting a NIOSH certificate of approval; (4) provides national recommendations for effective conformity assessment programs associated with non-respiratory PPT; (5) assesses research findings and translates them into effective conformity assessment recommendations for NIOSH policy, standards, regulations, and surveillance practices, for new protective technologies or special applications of existing technologies; (6) leads NIOSH participation in the development and promulgation of national and international consensus standards, conformity assessment program criteria and guidance, establishment of Federal regulations where necessary, and assesses of economic impact of Federal regulations; (7) prepares criteria for proper selection, recommends national guidance for effective use (*e.g.* cautions, limitations, and restrictions of use) and maintenance, and provides technical support; (8) plans and conducts public meetings to solicit or provide information concerning technology and conformity assessment practices; and (9) prepares and disseminates national reports related to conformity assessment of PPT.

Evaluation and Testing Branch (CCLH). (1) Conducts evaluations and tests in accordance with prescribed standard test procedures of RPD in support of NIOSH conformity assessment functions that lead to a NIOSH certificate of approval or its revocation; (2) conducts quality management system in-plant manufacturing-site evaluations including post market surveillance, and documents finding and recommendations in proper reports; (3) conducts evaluation and testing of PPT for various purposes, and prepares reports for dissemination to the public; (4) provides testing support to the NPPTL research and standards development initiatives; (5) develops evaluation methodologies, and unique test procedures to address new protective technologies or special applications of existing technologies; (6) conducts post market evaluations of NIOSH-certified RPD including the long-term field evaluation program, and prepares technical information and reports to improve standards for certification, selection, care, and use; (7) administers and conducts surveillance of field deployed PPT to evaluate conformance to applicable regulation, consensus standards, and NIOSH policy; (8) conducts investigations of PPT associated with complaints of nonconformance and/or concerns related to adverse health and safety including evaluations and analysis associated with NIOSH-certified respirators (*e.g.* certified product investigation process), and evaluating respirators and protective clothing submitted in conjunction with the NIOSH Fire Fighter Fatality Investigation and Prevention Program investigations conducted by the Division of Safety Research ; and (9) maintains and improves laboratory capabilities to perform evaluation and testing of PPT including innovative technologies, implements a laboratory quality program (*e.g.,* ISO 17025) to ensure quality and continuous improvement of PPT evaluations and tests, administers and maintains a chain of custody program to secure technologies or products obtained for evaluation and testing, and conducts an internal audit function to assure evaluation and testing are carried out in accordance policy and standard procedures.

James Seligman,

Acting Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2015–14686 Filed 6–15–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting of the aforementioned committee:

Time and Date:

1:30 p.m.–2:30 p.m. (EDT), July 17, 2015

Place: This meeting will be held by teleconference. To participate in the teleconference, please dial (877) 930–8819 and enter code 1579739.

Status: Open to the public, limited only by the availability of telephone ports. The public is welcome to participate during the public comment period, tentatively scheduled from 2:20 p.m. until 2:25 p.m.

Purpose: The Advisory Committee to the Director, CDC, shall advise the Secretary, HHS, and the Director, CDC, on policy and broad strategies that will enable CDC to fulfill its mission of protecting health through health promotion, prevention, and preparedness. The committee recommends ways to prioritize CDC's activities, improve results, and address health disparities. It also provides guidance to help CDC work more effectively with its various private and public sector constituents to make health protection a practical reality.

Matters for Discussion: The Advisory Committee to the Director will receive an update from the External Laboratory Safety Workgroup.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Carmen Villar, MSW, Designated Federal Officer, ACD, CDC, 1600 Clifton Road NE., M/S D–14, Atlanta, Georgia 30333;

Telephone (404) 639–7158; Email: *GHickman@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Claudette Grant,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2015–14749 Filed 6–15–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Applications for New Awards; Independent Living Administration

AGENCY: Administration for Community Living, Department of Health and Human Services.

ACTION: Notice.

Overview Information:

Independent Living Administration—Centers for Independent Living.

Notice inviting applications for new awards for fiscal year (FY) 2015.

Catalog of Federal Domestic Assistance (CFDA) Number: 93.432.

Note: This notice invites applications for separate competitions. For funding and other key information for this competition, see the chart in the *Award Information* section of this notice.

DATES: Applications Available: June 16, 2015.

Note: On July 22, 2014, President Obama signed the Workforce Innovation Opportunity Act (WIOA). WIOA was effective immediately. One provision of WIOA transferred the Centers for Independent Living (CIL) program from the Department of Education to the Administration for Community Living (ACL) in the Department of Health and Human Services. In addition, the CIL program will be placed in Independent Living Administration (ILA) within ACL. For FY 2015, all CIL program notices will be published as ACL notices, and ACL will make all CIL awards. ILA will post previously-approved application kits to grants.gov, and CIL applications submitted to grants.gov.

Date of Pre-Application Meeting: July 7, 2015.

Deadline for Notice of Intent to Apply: July 21, 2015.

Deadline for Transmittal of Applications: August 17, 2015.

Full Text of Announcement

I. Funding Opportunity Description

Purpose of Program: The purpose of the Center for Independent Living program provides support for planning, conducting, administering, and evaluating centers for independent living (centers) that comply with the standards and assurances in section 725 of part C of title VII of the Rehabilitation Act of 1973, as amended by the Workforce Innovation and Opportunity Act (WIOA, Pub. L. 113–128, consistent with the design included in the State plan for establishing a statewide network of centers.

Program Authority: 29 U.S.C. 796f–1.

Applicable Regulations: (a) The Department of Health and Human Services General Administrative Regulations in 45 CFR part 75 (b) Audit Requirements for Federal Awards in 45 CFR part 75 Subpart F; (c) 45 CFR part 75 Non-procurement Debarment and Suspension; (d) 45 CFR part 75 Requirement for Drug-Free Workplace (Financial Assistance); The regulations for this program in 45 CFR part 350.

Note: The regulations in 34 CFR part 79 apply to all applicants except federally recognized Indian tribes.

II. Award Information

Type of Award: Discretionary grant.

Estimated Available Funds: $249,142.

Estimated Number of Awards: 2.

States and outlying areas	Estimated available funds	Estimated number of awards
American Samoa	$154,046	1
Guam	95,096	1

Note: The Department is not bound by any estimates in this notice.

Project Period: Up to 60 months.

III. Eligibility Information

1. *Eligible Applicants:* To be eligible for funding, an applicant must—

(a) Be a consumer-controlled, community-based, cross-disability, nonresidential, private nonprofit agency;

(b) Have the power and authority to—

(1) Carry out the purpose of part C of title VII of the Act and perform the functions listed in section 725(b) and (c) of the Act and subparts F and G of 34 CFR part 366 within a community located within a State or in a bordering State; and

(2) Receive and administer—

(i) Funds under 34 CFR part 366;

(ii) Funds and contributions from private or public sources that may be used in support of a center; and

(iii) Funds from other public and private programs;

(c) Be able to plan, conduct, administer, and evaluate a center consistent with the standards and assurances in section 725(b) and (c) of the Act and subparts F and G of 34 CFR part 366;

(d) Either—

(1) Not currently be receiving funds under part C of chapter 1 of title VII of the Act; or

(2) Propose the expansion of an existing center through the establishment of a separate and complete center (except that the governing board of the existing center may serve as the governing board of the new center) at a different geographical location;

(e) Propose to serve one or more of the geographic areas that are identified as unserved or underserved by the States and Outlying Areas listed under *Estimated Number of Awards;* and

(f) Submit appropriate documentation demonstrating that the establishment of a new center is consistent with the design for establishing a statewide network of centers in the State plan of the State or Outlying Area whose geographic area or areas the applicant proposes to serve.

2. *Cost Sharing or Matching:* This competition does not require cost sharing or matching.

IV. Application and Submission Information

1. *Address to Request Application Package:* You can obtain an application package via grants.gov, or by contacting Veronica Hogan: U.S. Department of Health and Human Services, 400 Maryland Avenue SW., Room 5044, PCP, Washington, DC 20202–2800.