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) ways to enhance the quality, utility, and clarity of the information to be collected;(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; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information. Written comments should be received within 60 days of this notice.

Proposed Project

National Occupational Research Agenda (NORA) 2016 Decade Review—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The National Institute for Occupational Safety and Health (NIOSH) is responsible for conducting research and making recommendations to prevent worker injury and illness, as authorized in Section 20(a)(1) of the Occupational Safety and Health Act (29 U.S.C. 669). In 1995–6, NIOSH saw an opportunity to enhance its ability to accomplish its mission through partnerships that involved a broad national stakeholder base in occupational safety and health. With stakeholder input, NIOSH developed and launched a decade-long partnership program titled the National Occupational Research Agenda (NORA) in 1996. Participation in NORA includes stakeholders from universities, large and small businesses, professional societies, government agencies, and worker organizations. After an internal management review of the first decade of NORA, conducted in 2005, NIOSH launched the second decade of NORA (2006–2016) structured for even greater national impact. This information collection is a necessary part of a larger internal NIOSH management review of the second decade of NORA. The results of this review will inform NIOSH decisions about how to structure a third decade of NORA (2016–2026) for maximum effectiveness and impact.

The second decade of NORA was based on a new sector structure to better move research to practice within workplaces. The work of the sectors is managed through a partnership structure of sector councils. Each council develops and maintains an agenda for the decade for its sector. The sector agendas become part of the national agenda for improvements in occupational safety and health through research and partnerships. Representing all stakeholders, the councils use an open process to set goals, develop strategies, encourage partnerships, and promote improved workplace practices.

NIOSH is requesting a 12-month OMB approval to administer a survey to NORA council members and leaders. The collection of information is necessary for NIOSH management to assess the efficiency and effectiveness of the NORA sector councils. The target population is all current and former members and leaders of each of the ten NORA Sector Councils. The web-based questionnaire requests information on satisfaction with the efficiency of the council and its processes, on impacts made in the sector during the second decade, and suggestions for improving the effectiveness and impact of NORA in the future. Without this data collection, NIOSH's internal management review of NORA would lack critical stakeholder input from its many non-Federal partners.

A 16-item questionnaire has been developed and will be sent to all 352 non-Federal NORA Sector council members or leaders. A pilot test of the questionnaire was conducted by asking eight NIOSH employees who are a leader of a NORA sector council to complete the questionnaire and provide feedback. Respondents to the pilot test estimated the questionnaire requires approximately 15 minutes to complete. The total estimated burden is 88 hours. There is no cost to respondents other than their time.

Estimated Annualized Burden Hours

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)	Total burden (in hrs.)
Council member or leader	Council Questionnaire	352	1	15/60	88
Total					88

Leroy Richardson,

Chief, Information Collection Review, Office Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2014–12114 Filed 5–23–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; Extension of Certification of Maintenance of Effort on Help America Vote Act

AGENCY: Administration on Intellectual & Developmental Disabilities, Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the 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.

DATES: Submit written or electronic comments on the collection of information by July 28, 2014.

ADDRESSES: Submit electronic comments on the collection of information to: *clare.barnett@acl.hhs.gov*.

Submit written comments on the collection of information to Administration for Community Living, One Massachusetts Avenue NW., Washington, DC 20201, attention Clare Barnett.

FOR FURTHER INFORMATION CONTACT: Clare Barnett, Program Specialist, Administration for Community Living, One Massachusetts Avenue NW., Washington, DC 20201.

SUPPLEMENTARY INFORMATION: This notice solicits comments on the information collection requirements relating to the Help America Vote Act (HAVA)), Public Law 107–252, Title II, Subtitle D, Part 2, Sections 261 to 265 (HAVA Narrative Annual Report). 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 request 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 notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on: (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; (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.

The Help America Vote Act (HAVA) Narrative Report from States and Units of Local Government is required by federal statute and regulation, the Help America Vote Act HAVA), Public Law 107–252, Title II, Subtitle D, Part 2, Sections 261 to 265, Payments to States and Units of Local Government to Assure Access for Individuals with Disabilities (42 U.S.C. 15421–25). The report is provided in writing to the Administration for Community Living, Administration on Intellectual and Developmental Disabilities. Each State or Unit of Local Government must prepare and submit an annual report at the end of every fiscal year. The report addresses the activities conducted with the funds provided during the year. The information collected from the annual report will be aggregated into an annual profile of how States have utilized the funds and establish best practices for election officials. It will also provide an overview of the State election goals and accomplishments and permit the Administration on Intellectual & Developmental Disabilities to track voting progress to monitor grant activities. ACL estimates the burden of this collection of information as follows: 55 Chief Election officials respond annually which should be an average burden of 20 hours per State per year or a total of 1,100 hours for all states annually.

Dated: May 15, 2014.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2014–11989 Filed 5–23–14; 8:45 am]

BILLING CODE4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Allergy, Immunology, and Transplantation Research.

Date: June 17, 2014.

Time: 10:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health Room 3147, 6700B Rockledge Drive Bethesda, MD 20817 (Telephone Conference Call).

Contact Person: James T. Snyder, Ph.D., Scientific Review Officer Scientific Review Program, Division of Extramural Activities/NIAID National Institutes of Health, 6700B Rockledge Drive, MSC 7616 Bethesda, MD 20892–7616, 301–435–1614, *james.snyder@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: May 20, 2014.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014–12081 Filed 5–23–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Heart, Lung, and Blood Initial Review Group; Clinical Trials Review Committee.

Date: June 23–24, 2014.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Keary A Cope, Ph.D. Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room