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