FOR FURTHER INFORMATION CONTACT: Kelly Powell, HR Specialist, at 202–942–1681.

SUPPLEMENTARY INFORMATION: Title 5, U.S. Code, 4314(c)(4), requires that the appointment of Performance Review Board members be published in the **Federal Register** before Board service commences. The following persons will serve on the Federal Retirement Thrift Investment Board's Performance Review Boards which will review initial summary ratings to ensure the ratings are consistent with established performance requirements, reflect meaningful distinctions among senior executives based on their relative performance and organizational results and provide recommendations for ratings, awards, and pay adjustments in a fair and equitable manner: James Petrick, Renee Wilder, and Karen Vaughn Peck.

Megan Grumbine,

Deputy General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 2015–28735 Filed 11–10–15; 8:45 am]

BILLING CODE 6760–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; Extension State Plan for Independent Living (SPIL) Public Law (105–220) for the State Independent Living (SILS) and Centers for Independent Living (CIL) Program Authorized by Title VII, Chapter 1, of the, as Amended by the Workforce Innovation and Opportunity Act (WIOA, Pub. L. 113–128) [Rehabilitation Act]

AGENCY: Center for Independent Living Administration, Administration for Community Living (ACL), HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL), Independent Living Administration is announcing an opportunity for public comment on the proposed collection of certain information. 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. This notice solicits comments on the information collection requirements relating to the State Plan for Independent Living (SPIL).

DATES: Submit written or electronic comments on the collection of information by January 11, 2016.

ADDRESSES: Submit electronic comments on the collection of information to: *veronica.hogan@acl.hhs.gov.* Submit written comments on the collection of information to Administration for Community, Independent Living Administration, 550 12th Street Southwest, PCP Building, Room 5044, Washington, DC 20202, attention Veronica Hogan.

FOR FURTHER INFORMATION CONTACT: Veronica Hogan, Grant Management Specialist, (202) 245–7378 or by email *veronica.hogan@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 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/ILA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL/ILA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of ACL/ILA's functions, including whether the information will have practical utility; (2) the accuracy of ACL/ILA 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 Independent Living Program is required by federal statute and regulation requires the collection of this information every three years. The three-year period for the next SPIL is FY 2018–2020. The SPIL provided in writing to the Administration for Community Living, Administration on Disabilities, Independent Living Administration. The five core services are: Advocacy, information and referral, independent living skills training, peer counseling, and transition services. WIOA included three prongs to the 5th core service:

- Facilitating the transition of individuals with significant disabilities from nursing homes and other institutions to home and community-based residences, with the requisite supports and services;
- Provide assistance to individuals with significant disabilities who are at risk of entering institutions so that the individuals may remain in the community, and
- Facilitate the transition of youth who are individuals with significant disabilities, who were eligible for individualized education programs under section 614(d) of the Individuals with Disabilities Act (20 U.S.C. 1414(d)), and who have completed their secondary education or otherwise left school, to postsecondary life.

ACL estimates the burden of this collection of information as follows: 56 SPIL respond annually which should be an average burden of 60 hours for each grantee. The aggregate hour burden for all grantees is an estimated 3,360 hours (56 grantees × 60 hours each). These estimated hours include the time required for reading, studying and planning for the new SPIL; conducting required public hearings; gathering and reviewing pertinent information; completing the SPIL assurances and narrative sections; reviewing draft and final versions of the completed SPIL; and submission of the final SPIL to ACL.

Dated: November 5, 2015.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2015–28745 Filed 11–10–15; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; Semi-Annual and Final Reporting Requirements for Discretionary Grant Programs

AGENCY: 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. This notice solicits comments on the information collection requirements relating to the continuation of an existing collection for Performance Progress Reports previously approved for discretionary grants funded by the U.S. Administration for Community Living (ACL).

DATES: Submit written or electronic comments on the collection of information by January 11, 2016.

ADDRESSES: Submit electronic comments on the collection of information to: *lori.stalbaum@acl.hhs.gov.* Submit written comments on the collection of information to Lori Stalbaum, Administration on Community Living, Washington, DC 20201 or by fax to Lori Stalbaum at 202–357–3469.

FOR FURTHER INFORMATION CONTACT: Lori Stalbaum at 202–357–3452 or *lori.stalbaum@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 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 Administration for Community Living (ACL) plans to continue an existing approved collection of information for semi-annual and final reports pursuant to the requirements of its discretionary grant programs. Through its discretionary grant programs, ACL supports projects for the purpose of developing and testing new knowledge and program innovations with the potential for contributing to the independence, well-being, and health of older adults, people with disabilities across the lifespan, and their families and caregivers. Deliverables required by ACL of all grantees are semi-annual and final reports, as provided for in the Department of Health and Human Services regulations, 45 CFR part 74, Section 74.51. These grantee performance reporting requirements can be found on AoA's Web site at *http://www.acl.gov/Funding_Opportunities/Grantee_Info/Reporting.aspx.* ACL estimates the burden of this collection of information as follows: *Frequency:* Semi-annually with the Final report taking the place of the semi-annual report at the end of the final year of the grant. *Respondents:* States, public agencies, private nonprofit agencies, institutions of higher education, and organizations including tribal organizations. *Estimated Number of Responses:* 600. *Total Estimated Burden Hours:* 12,000.

Dated: November 5, 2015.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2015–28744 Filed 11–10–15; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–N–0001]

Endocrinologic and Metabolic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Endocrinologic and Metabolic Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on December 14, 2015, from 8 a.m. to 5 p.m.

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993–0002. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

Contact Person: Stephanie L. Begansky, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, (301) 796–9001, FAX: (301) 847–8533, *EMDAC@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: The committee will discuss the results of the IMProved Reduction of Outcomes: Vytorin Efficacy International Trial (IMPROVE–IT). IMPROVE–IT was a clinical trial that studied the effect of ezetimibe/simvastatin compared with simvastatin on the occurrence of cardiovascular events in patients with recent acute coronary syndrome. The results from this trial have been submitted to support supplemental new drug applications 21445/S–038 and 21687/S–054, ZETIA (ezetimibe) and VYTORIN (ezetimibe/simvastatin) tablets, respectively, by MSD International GmbH. The proposed indication for ZETIA (in combination with a statin) and VYTORIN is to reduce