Place: National Institutes of Health, Room 5F100, 5601 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: Andrea L. Wurster, Ph.D., Scientific Review Officer Scientific Review Program, Division of Extramural Activities, Room 3G33B National Institutes of Health, NIAID, 5601 Fishers Lane, MSC 9823, Bethesda, MD 20899823, (240) 669–5062, *wurstera@mail.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: March 22, 2016.

Natasha M. Copeland,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–06871 Filed 3–25–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; 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: National Institute on Aging Special Emphasis Panel: Vascular Dysfunction in AD and Genetic Risk Factors.

Date: May 6, 2016.

Time: 11:30 a.m. to 3:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute on Aging, Gateway Building, Suite 2C212, 7201 Wisconsin Avenue, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Alexander Parsadanian, Ph.D., Scientific Review Officer, National Institute on Aging, Gateway Building 2C/212, 7201 Wisconsin Avenue, Bethesda, MD 20892, 301–496–9666, PARSADANIANA@NIA.NIH.GOV.

Name of Committee: National Institute on Aging Special Emphasis Panel: 2016 Beeson Review

Date: May 26, 2016

Time: 8:00 a.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications

Place: DoubleTree by Hilton, 8120 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Alexander Parsadanian, Ph.D., Scientific Review Office, National Institute on Aging, Gateway Building 2C/212, 7201 Wisconsin Avenue, Bethesda, MD 20892, 301–496–9666, PARSADANIANA@NIA.NIH.GOV.

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, Aging Research, National Institutes of Health, HHS).

Dated: March 22, 2016.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–06870 Filed 3–25–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; U.S. Nuclear Medicine Technologists Study (NCI)

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute, the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; The quality, utility, and clarity of the information to be collected; and minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

To Submit Comments and For Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact*: Michele M. Doody, Radiation Epidemiology Branch, National Cancer Institute, 9609 Medical Center Drive, Room 7E566, Rockville, MD 20850, or call non-toll-free at 301–414–0308. Or Email your request, including your address to: *doodym@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Proposed Collection: US Nuclear Medicine Technologists Study, 0925–0656, Expiration Date 04/30/2015—REINSTATEMENT WITH CHANGE, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: We propose to collect, from U.S. nuclear medicine technologists (USNMT) certified after 1980, historical information about nuclear medicine procedures performed, radioisotopes used, related work and safety practices, and places of employment. The primary objectives of the current feasibility effort are: (a) To identify a cohort of nuclear medicine technologists certified after 1980 by the American Registry of Radiologic Technologists (ARRT) and/or the Nuclear Medicine Technologist Certification Board (NMTCB); and (b) to characterize individual organ-specific occupational radiation doses from radioisotope procedures. More recently certified technologists, who specialized in nuclear medicine, are expected to have greater exposures to radioisotopes than the general radiologic technologists in the U.S. Radiologic Technologist (USRT) cohort owing to performing such procedures with greater frequency. The proposed USNMT study would be a direct follow-on to the USRT Study to assess health risks associated with occupational exposure to these much higher-energy radiopharmaceuticals.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 125.