information collection provisions in the guidance.

Description of Respondents: The likely respondents include businesses engaged in the manufacture or sale of food, food ingredients, and substances

used in materials that come into contact with food.

In the **Federal Register** of August 25, 2016 (81 FR 58517), FDA published a 60-day notice requesting public comment on the proposed collection of

information. No comments were received.

We estimate the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN 1

21 CFR section	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
25.15 (a) & (d) (to cover CEs under 25.32(i))	47 1 3	1 1 1	47 1 3	8 8 8	376 8 24
25.40 (a) & (c) EAs	57	1	57	180	10,260

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

The estimates for respondents and numbers of responses are based on the annualized numbers of petitions and notifications qualifying for categorical exclusions listed under § 25.32(i) and (q) that the Agency has received in the past 3 years. Please note that, in the past 3 years, there have been no submissions that requested an action that would have been subject to the categorical exclusion in § 25.32(o). To avoid counting this burden as zero, we have estimated the burden for this categorical exclusion at one respondent making one submission a year for a total of one annual submission. The burden for submitting a categorical exclusion is captured under § 25.15(a) and (d).

To calculate the estimate for the hours per response values, we assumed that the information requested in this guidance for each of these three categorical exclusions is readily available to the submitter. For the information requested for the exclusion in § 25.32(i), we expect that submitter will need to gather information from appropriate persons in the submitter's company and prepare this information for attachment to the claim for categorical exclusion. We believe that this effort should take no longer than 8 hours per submission. For the information requested for the categorical exclusions in § 25.32(o) and (q), the submitters will almost always merely need to copy existing documentation and attach it to the claim for categorical exclusion. We believe that collecting this information should also take no longer than 8 hours per submission.

For the information requested for the environmental assessments in § 25.40(a) and (c), we believe that submitters will submit an average of 57 environmental assessments annually. We estimate that each submitter will prepare an EA within 3 weeks (120 hours) and revise

the EA based on Agency comments (between 40 to 60 hours), for a total preparation time of 180 hours. The burden relating to this collection has been previously approved under OMB control number 0910–0322,

"Environmental Impact Consideration— 21 CFR part 25". Upon approval of this collection of information by OMB, FDA will revise OMB control number 0910-0322 to remove the annual reporting burden for categorical exclusions and environmental assessment requests related to food additive petitions, color additive petitions, requests for exemption from regulation as a food additive, and submission of a food contact notification for a food contact substance. The future burden for categorical exclusion or environmental assessments for these requests will be captured under OMB control number 0910-0541, this collection of information.

Dated: November 15, 2016.

Leslie Kux,

Associate Commissioner for Policy.
[FR Doc. 2016–27943 Filed 11–18–16; 8:45 am]
BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Privacy Act of 1974; System of Records Notice

AGENCY: Department of Health and Human Services (HHS), Office of the Secretary (OS)

ACTION: Notice to establish a new system of records, and to delete related systems.

SUMMARY: In accordance with the requirements of the Privacy Act of 1974, as amended, HHS is establishing a new,

department-wide system of records, System No. 09-90-1601 "Outside Experts Recruited for Non-FACA Activities," and deleting four related systems of records that are obsolete or that will be rendered duplicative by the new system. The new system will cover recruitment and other administrative records about individuals outside the HHS workforce who serve or are considered for service on HHS missionrelated committees and other assignments requiring specific outside expertise or experience (excluding those that are subject to the Federal Advisory Committee Act (FACA), which are covered under System No. 09–90–0059). The new department-wide System No. 09-90-1601 and the related system deletions are more fully explained in the **SUPPLEMENTARY INFORMATION** section of this Notice.

DATES: The new system of records established in this Notice is effective upon publication, with the exception of the routine uses. The routine uses will be effective 30 days after publication of this Notice, unless comments are received that warrant a revision to this Notice. Written comments on the Notice should be submitted within 30 days. The deletion of System Numbers 09–20–0168, 09–30–0049, 09–37–0022, and 09–90–0080 will be effective 30 days after publication of this Notice.

ADDRESSES: The public should address written comments to: Beth Kramer, HHS Privacy Act Officer, FOIA/PA Division, Hubert H. Humphrey Building—Suite 729H, 200 Independence Avenue SW., Washington, DC 20201, beth.kramer@hhs.gov.

FOR FURTHER INFORMATION CONTACT: Beth Kramer, HHS Privacy Act Officer, FOIA/ PA Division, Hubert H. Humphrey Building—Suite 729H, 200 Independence Avenue SW., Washington, DC 20201, beth.kramer@hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Explanation of New System No. 09–90–1601

The records to be covered in the new system of records are similar in type and function to the records covered in System No. 09-90-0059, which pertain to individuals who serve or are considered for service on committees that are subject to the Federal Advisory Committee Act (FACA), 5 U.S.C. App., et seq.; the key difference is that they will be about outside individuals serving or considered for service on mission-related committees and other activities that are not subject to FACA. Following are the non-FACA-related programs at HHS that recruit and utilize individuals with outside expertise or experience and maintain records about the outside individuals in systems that retrieve the records by personal identifier:

- Curricula Vitae of Consultants to the National Center for Health Statistics (NCHS) within the Centers for Disease Control and Prevention (CDC) (formerly covered under SORN 09-20-0168). This program maintains records about individuals with special expertise, training, and professional experience who may be enlisted to assist CDC/ NCHS as consultants. The records are used by CDC/NCHS to select individuals to participate in assignments such as: planning and conducting surveys, studies, statistical reporting programs, and statistical analyses of data; providing training and technical assistance; and planning and conducting conferences. These records currently are covered under SORN No. 09-20-0168, which is being deleted and subsumed under the new departmentwide SORN No. 09-90-1601.
- The Food and Drug Administration (FDA) Patient Representative Program. This program enlists individuals with patient advocacy experience to serve as patient representatives on both FACA committees and non-FACA assignments. For example, patient representatives may provide input that is used in making decisions to approve devices or drugs, or may contribute to discussions at presentations and conferences. Records about patient representatives are retrieved by the representatives' names, and will be covered under either SORN No. 09–90–0059 or the new department-wide SORN No. 09-90-1601, depending on whether the records pertain to service on a FACA committee or service on a non-FACA assignment.

- Peer Review Programs at the Administration for Children and Families (ACF), Health Resources and Services Administration (HRSA), and Substance Abuse and Mental Health Services Administration (SAMHSA) that recruit and use outside individuals to serve on peer review committees formed to review applications for grants and cooperative agreements. These programs exist in several HHS components, but only ACF, HRSA, and SAMHSA sometimes use a personal identifier (i.e., name) to retrieve administrative records about the outside individuals they recruit and use. Other components (including the Office of the Assistant Secretary for Health (OASH), Centers for Medicare & Medicaid Services (CMS), and National Institutes of Health (NIH)) use only non-personal identifiers (e.g., expertise type, or funding opportunity announcement number) for retrieval.
- Consultants on Other SAMHSA
 Projects (formerly covered under SORN
 09–30–0049). SAMHSA contractors
 arrange for outside consultants to be
 used in other SAMHSA programs
 (besides peer review programs) when
 technical assistance is needed in
 conferences, meetings, and evaluation
 projects that involve a specialized area
 of research, review, or advice.

A report on the new system of records has been sent to Congress and OMB in accordance with 5 U.S.C. 552a(r).

II. Deletion of Four Related Systems of Records

The following systems of records are being deleted as duplicative of new department-wide System No. 09–90–1601:

- 09–20–0168 Curricula Vitae of Consultants to the National Center for Health Statistics
- 09–30–0049 Consultant Records Maintained by SAMHSA Contractors

The following system of records is being deleted as duplicative of System No. 09–90–0059 Federal Advisory Committee Membership Files as to files that pertain to candidates for FACA committees, and as duplicative of new department-wide System No. 09–90–1601 as to files that pertain to candidates for non-FACA committees and other activities:

- 09–90–0080 The Secretary's Advisory Committee Candidate Files
- The following system of records is being deleted because it is obsolete and the records no longer exist:
- 9-37-0022 Records of Health Experts Maintained by the Office of International Health

III. The Privacy Act

The Privacy Act (5 U.S.C. 552a) governs the means by which the U.S. Government collects, maintains, and uses information about individuals in a system of records. A "system of records" is a group of any records under the control of a federal agency from which information about an individual is retrieved by the individual's name or other personal identifier. The Privacy Act requires each agency to publish in the Federal Register a system of records notice (SORN) identifying and describing each system of records the agency maintains, including the purposes for which the agency uses information about individuals in the system, the routine uses for which the agency discloses such information outside the agency, and how individual record subjects can exercise their rights under the Privacy Act.

Dated: November 1, 2016.

Beth Kramer,

Privacy Act Officer, FOIA/Privacy Act Division, Assistant Secretary for Public Affairs, Department of Health and Human Services.

Notice of Deletion of Related Systems

The following systems of record are deleted, effective 30 days after publication of this Notice:

- 1. 09–20–0168 Curricula Vitae of Consultants to the National Center for Health Statistics
- 2. 09–30–0049 Consultant Records Maintained by SAMHSA Contractors
- 3. 09–90–0080 The Secretary's Advisory Committee Candidate Files
- 4. 09–37–0022 Records of Health Experts Maintained by the Office of International Health

SYSTEM NUMBER:

09-90-1601

SYSTEM NAME:

Outside Experts Recruited for Non-FACA Activities

SECURITY CLASSIFICATION:

Unclassified.

SYSTEM LOCATION:

Physical locations include:

- CDC program offices that recruit consultants to assist in statistical projects and reporting programs conducted or sponsored by NCHS, in Atlanta, GA and Hyattsville, MD;
- FDA's committee management office in Silver Spring, MD;
- Program offices at ACF in Washington, DC, at HRSA in Rockville, MD, and at SAMHSA in Rockville, MD,

that recruit individuals to serve as peer reviewers; and

• Locations of SAMHSA contractors that arrange use of consultants on SAMHSA projects.

CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:

Records in this system pertain to individuals outside the HHS workforce who serve or are considered for service on HHS mission-related committees or other assignments that require specific outside expertise or experience (for example, medical, scientific, or manufacturing expertise, or patient advocacy experience), but that are not subject to the Federal Advisory Committee Act (FACA), 5 U.S.C. App., et seq.

CATEGORIES OF RECORDS IN THE SYSTEM:

The records consist of recruitment and other administrative records, including:

- An application and resume or curricula vitae, describing the individual's qualifications;
- Nomination/recommendation records, or other records used in evaluating an individual's qualifications and any potential conflicts of interest and selecting an individual for a specific assignment; and
- Records used to plan and arrange the individual's participation in the assigned activities, including scheduling records and records used to coordinate parking, badging, and payment of any stipend or honorarium.

The records may contain these data elements:

- The individual's name and other identifying information (*e.g.*, sex, place and date of birth);
- Contact information (e.g., home and business addresses, telephone numbers, email addresses);
- Occupation, job titles, employers, employment status and history, and whether currently employed by the federal government;
- Work and organizational affiliations, memberships, credentials, and licenses;
- Degrees held, and general educational and/or experience background;
- Racial classification or ethnic background;
- Areas of specialization, expertise, or experience, and special qualifications (e.g., language or technical skills, ability to drive to an assignment);
- Dates and descriptions of past assignments or past experience;
- Sources and references, and any information provided by sources/ references; and

• Information about availability and any special needs.

Any special needs, medical condition, or similar information contained in an individual's records is maintained and used in accordance with relevant provisions of the Rehabilitation Act of 1973, as amended, 29 U.S.C. 791 et seq., and implementing regulations at 29 CFR parts 1614 and 1630, and the Genetic Information Nondiscrimination Act of 2008 at 42 U.S.C. 2000ff et seq.

AUTHORITY FOR MAINTENANCE OF THE SYSTEM:

For CDC/NCHS Consultant Records: 42 U.S.C. 242b(b)(3).

For FDA Patient Representative Records: 21 U.S.C. 360bbb–8c, 371 et seq., 379d–1(b)(1)(A).

For ACF Peer Reviewer Records: 42 U.S.C. 799(f), 806(e).

For HRSA Peer Reviewer Records: 42 U.S.C. 799(f), 806(e).

For SAMHSA Peer Reviewer and Other Consultant Records: 42 U.S.C. 241, 249(c), 290aa et seq., 290aa–5, 290bb et seq., 290bb–21 et seq., 290bb– 31 et seq., 5121 et seq., 10801 et seq.; 8 U.S.C. 1522 note; Executive Order 12341.

See also: 5 U.S.C. 3109.

PURPOSE(S):

The records will be used within the agency on a need-to-know basis for the purpose of staffing committees and other assignments and managing administrative matters pertaining to individuals serving on committees and other assignments, including to:

- Prepare reports and lists of past, present, and recommended members, vacancies, acceptances, and separations;
- Send recruitment notices to individual prospective candidates, and send informational notices to selectees;
- Identify qualified candidates and document the selections; and
- Manage and coordinate the selected individuals' participation in assignment activities (including sharing information within the agency to coordinate aspects such as badging, parking, travel, training, and payment of any stipend or honorarium).

ROUTINE USES OF RECORDS MAINTAINED IN THE SYSTEM, INCLUDING CATEGORIES OF USERS AND THE PURPOSES OF SUCH USES:

In addition to the statutory disclosures of information permitted in the Privacy Act at 5 U.S.C. 552a(b)(2) and (b)(4)–(11), HHS may make the following disclosures of information about an individual from this system of records to parties outside the agency without the individual's prior, written consent:

1. Disclosures may be made to federal agencies and Department contractors

that have been engaged by HHS to assist in accomplishment of an HHS function relating to the purposes of this system of records and that have a need to have access to the records in order to assist HHS in performing the activity. Any contractor will be required to comply with the requirements of the Privacy Act.

- 2. Records may be disclosed to parties such as educational institutions, current and former employers, and qualified experts, when necessary to check or obtain an opinion about a candidate's qualifications.
- 3. Records about consultants and patient advocates may be disclosed to parties organizing or hosting assignment activities, such as grantee institutions and federal, foreign, state, tribal, local, and other government agencies and public authorities (e.g., U.S. Embassies and Ministries of Health), when necessary to apprise them of an individual's qualifications for the assignment or coordinate the individual's participation in the activities.
- 4. Records may be disclosed to supervisors and administrative assistants at the individual's place of employment, for administrative purposes such as coordinating the individual's participation in the activities.
- 5. Records may be disclosed to external parties that audit committee or assignment activities.
- 6. Relevant information will be included in any required reports to the President, the Office of Management and Budget (OMB), and the General Services Administration (GSA) about committees and other assignments that are mission-related.
- 7. Information may be disclosed to the U.S. Department of Justice (DOJ) or to a court or other tribunal, when:
- a. The agency or any component thereof, or
- b. Any employee of the agency in his or her official capacity, or
- c. Any employee of the agency in his or her individual capacity where DOJ has agreed to represent the employee, or d. The United States Government,
- is a party to litigation or has an interest in such litigation and, by careful review, HHS determines that the records are both relevant and necessary to the litigation and that, therefore, the use of such records by the DOJ, court or other tribunal is deemed by HHS to be compatible with the purpose for which the agency collected the records.
- 8. Records may be disclosed to student volunteers and other individuals performing functions for the

Department but technically not having the status of agency employees, if they need access to the records in order to perform their assigned agency functions.

9. Disclosures may be made to the National Archives and Records Administration (NARA) and/or the General Services Administration (GSA) for the purpose of records management inspections conducted under 44 U.S.C. 2904 and 2906.

10. Information may be disclosed to a Member of Congress or a Congressional staff member in response to a written inquiry of the Congressional office made at the written request of the constituent about whom the record is maintained. The Congressional office does not have any greater authority to obtain records than the individual would have if requesting the records directly.

11. Records may be disclosed to the U.S. Department of Homeland Security (DHS) if captured in an intrusion detection system used by HHS and DHS pursuant to a DHS cybersecurity program that monitors Internet traffic to and from federal government computer networks to prevent a variety of types of cybersecurity incidents.

12. Disclosures may be made to appropriate federal agencies and Department contractors that have a need to know the information for the purpose of assisting the Department's efforts to respond to a suspected or confirmed breach of the security or confidentiality of information maintained in this system of records, when the information disclosed is relevant and necessary to that assistance.

POLICIES AND PRACTICES FOR STORING, RETRIEVING, ACCESSING, RETAINING, AND DISPOSING OF RECORDS IN THE SYSTEM—

STORAGE:

Records are stored in hard-copy files and electronic media.

RETRIEVABILITY:

Records are retrieved by the individual's name.

SAFEGUARDS:

Safeguards conform to the HHS
Information Security and Privacy
Program, http://www.hhs.gov/ocio/
securityprivacy/index.html. Information
is safeguarded in accordance with
applicable laws, rules and policies,
including the HHS Information
Technology Security Program
Handbook, all pertinent National
Institutes of Standards and Technology
(NIST) publications, and OMB Circular
A–130, Management of Federal
Resources. Records are protected from
unauthorized access through
appropriate administrative, physical,

and technical safeguards. These safeguards include protecting the facilities where records are stored or accessed with security guards, badges and cameras, securing hard-copy records in locked file cabinets, file rooms or offices during off-duty hours, limiting access to electronic databases to authorized users based on roles and two-factor authentication (user ID and password), using a secured operating system protected by encryption, firewalls, and intrusion detection systems, requiring encryption for records stored on removable media, and training personnel in Privacy Act and information security requirements. Records that are eligible for destruction are disposed of using destruction methods prescribed by NIST SP 800-88.

RETENTION AND DISPOSAL:

Records pertaining to recruitment and use of outside peer reviewers are destroyed three years after final action; they are retained longer if required for business use (see General Records Schedule (GRS) 1.2, Item 010, Grant and Cooperative Agreement Program Management Records). Records pertaining to recruitment and use of other outside individuals (e.g., experts, patient advocates, and members of mission-related non-FACA committees) are currently unscheduled. Unscheduled records must be retained indefinitely pending the agency's submission, and NARA's approval, of a disposition schedule. HHS anticipates proposing to NARA, as an appropriate retention period for these records, "three years after final action, or longer if required for business use" (similar to the period provided in GRS 1.2, Item 010) or "when no longer needed for administrative purposes" (similar to the periods applicable to similar records not retrieved by personal identifier which are not covered under this SORN; i.e.: N1-442-93-1, Item 37 for the Agency for Toxic Substances and Disease Registry's Curriculum Vitae Files, and NC1-235-82-1, Item 100-3 for the Office of the Secretary's Advisory Committee Candidate Resume Files).

SYSTEM MANAGER(S) AND ADDRESS(ES):

For CDC/NCHS Consultant Records:

• Centers for Disease Control and Prevention (CDC), Director, National Center for Health Statistics, OPHSS, Prince George's Metro IV Bldg., Rm. 7209, MS P08, Centers for Disease Control and Prevention, 3311 Toledo Road, Hyattsville, MD 20782

For FDA Patient Representative Records:

• Food and Drug Administration (FDA), Advisory Committee Oversight &

Management Staff, 10903 New Hampshire Avenue, Bldg. WO32, Rm. 5129, Silver Spring, MD 20993–002

For ACF Peer Reviewer Records:

• Administration for Children and Families (ACF), Privacy Act Contact, Office of Information Systems 330 C Street, NW., Washington, DC 20201

For HRSA Peer Reviewer Records:

• Health Resources and Services Administration (HRSA), Chief, Policy, Analysis & Training Branch, Division of Independent Review, Office of Federal Assistance Management, 5600 Fishers Lane, Rockville, MD 20857

For SAMHSA Peer Reviewer Records:

• Substance Abuse and Mental Health Services Administration (SAMHSA), Director, Division of Grant Review, 5600 Fishers lane, Rockville, MD 20852

For Other Consultant Records, Maintained by SAMHSA Contractors:

• Substance Abuse and Mental Health Services Administration (SAMHSA), Director, Division of Contracts Management, Office of Program Services, 5600 Fishers Lane, Rockville, MD 20852

NOTIFICATION PROCEDURE:

An individual who wishes to know if this system contains records about him or her should submit a written request to the relevant System Manager indicated above. The individual must verify his or her identity by providing either a notarization of the request or a written certification that the requester is who he or she claims to be and understands that the knowing and willful request for acquisition of a record pertaining to an individual under false pretenses is a criminal offense under the Privacy Act, subject to a five thousand dollar fine.

RECORD ACCESS PROCEDURE:

An individual seeking access to records about him in this system should submit a request following the same procedure indicated under "Notification Procedure."

CONTESTING RECORD PROCEDURE:

An individual seeking to amend the content of information about him or her in this system should contact the relevant System Manager indicated above and reasonably identify the record, specify the information contested, state the corrective action sought, and provide the reasons for the amendment, with supporting justification.

RECORD SOURCE CATEGORIES:

Most information is obtained directly from the individual record subject.

Information pertaining to references and recommendations is obtained from other private individuals, educational institutions, current and former employers, HHS program personnel, biographical reference books, private organizations, Members of Congress, and other government sources.

EXEMPTIONS CLAIMED FOR THE SYSTEM:

None.

[FR Doc. 2016–27959 Filed 11–18–16; 8:45 am] **BILLING CODE 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: National Heart, Lung, and Blood Institute Special Emphasis Panel; Human Virome in Heart, Lung, and Blood Health and Resilience.

Date: December 9, 2016.
Time: 8:30 a.m. to 5:00 p.m.
Agenda: To review and evaluate grant applications.

Place: Residence Inn Bethesda, 7335 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Kristen Page, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, 301–496–2434, kristen.page@nih.gov.

Name of Committee: National Heart, Lung, and Blood Institute Special Emphasis Panel; T32—Training Programs for Institutions that Promote Diversity.

Date: December 9, 2016.

Time: 11:00 a.m. to 12:00 p.m. Agenda: To review and evaluate g

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7189, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Stephanie L. Constant, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7189, Bethesda, MD 20892, 301– 443–8784, constantsl@nhlbi.nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: November 15, 2016.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–27876 Filed 11–18–16; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Drug Abuse; 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: National Institute on Drug Abuse Special Emphasis Panel, Multisite Clinical Trials SEP III.

Date: December 9, 2016.

Time: 1:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Telephone Conference Call).

Contact Person: Susan O. McGuire, Ph.D., Scientific Review Officer, Office of Extramural Affairs, National Institute on Drug Abuse, National Institutes of Health, DHHS, 6001 Executive Blvd., Room 4245, Rockville, MD 20852, (301) 827–5817, mcguireso@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos.: 93.279, Drug Abuse and Addiction Research Programs, National Institutes of Health, HHS)

Dated: November 15, 2016.

Natasha M. Copeland,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–27878 Filed 11–18–16; 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 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: Heart, Lung, and Blood Initial Review Group, NHLBI Institutional Training Mechanism Review Committee.

Date: December 9, 2016.

Time: 10:00 a.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7194, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Charles Joyce, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7194, Bethesda, MD 20892–7924, 301–435– 0288, cjoyce@nhlbi.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: November 15, 2016.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016-27875 Filed 11-18-16; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; 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