provides advice on priorities and activities related to the development, validation, scientific review, regulatory acceptance, implementation, and national and international harmonization of new, revised, and alternative toxicological test methods. Additional information about SACATM, including the charter, roster, and records of past meetings, can be found at <a href="http://ntp.niehs.nih.gov/go/167">http://ntp.niehs.nih.gov/go/167</a>.

Dated: May 4, 2010.

#### John R. Bucher,

Associate Director, National Toxicology Program.

[FR Doc. 2010–11318 Filed 5–11–10; 8:45 am]

BILLING CODE 4140-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# **Administration on Aging**

Agency Information Collection Activities; Proposed Collection; Comment Request; National Survey of Older Americans Act Title III Service Recipients

**AGENCY:** Administration on Aging, HHS. **ACTION:** Notice.

**SUMMARY:** The Administration on Aging (AoA) 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 information collection requirements contained in consumer assessment surveys that are used by AoA to measure program performance for programs funded under Title III of the Older Americans Act.

**DATES:** Submit written or electronic comments on the collection of information by July 12, 2010.

**ADDRESSES:** Submit electronic comments on the collection of information to:

valerie.cook@aoa.hhs.gov. Submit written comments on the collection of information to Valerie Cook, Administration on Aging, Office of Evaluation, Washington, DC 20201.

**FOR FURTHER INFORMATION CONTACT:** Valerie Cook 202–357–3583.

**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, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA'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 National Survey of Older Americans Act (OAA) Title III Service Recipients information collection, which builds on earlier national pilot studies and surveys, as well as performance measurement tools developed by AoA grantees in the Performance Outcomes Measures Project (POMP), will include consumer assessment surveys for the Congregate and Home-delivered meal nutrition programs; Case Management, Homemaker, and Transportation Services; and the National Family Caregiver Support Program. This information will be used by AoA to track performance outcome measures; support budget requests; comply with Government Performance and Results Act (GPRA) reporting requirements; provide national benchmark information for POMP grantees; and inform program development and management initiatives. Descriptions of previous National Surveys of OAA Participants can be found under the section on OAA Performance

Information on AoA's Web site at: http://www.aoa.gov/AoARoot/ Program Results/ OAA Performance.aspx. Copies of the survey instruments and data from previous National Surveys of OAA Participants can be found and queried using the AGing Integrated Database (AGID) at http://www.agidnet.org/. AoA estimates the burden of this collection of information as follows: Respondents: Individuals; Number of Respondents: 6,250; Number of Responses per Respondent: one; Average Burden per Response: 6000 at 30 minutes, 250 at 4 hours; Total Burden: 4,000.

Dated: May 6, 2010.

#### Kathy Greenlee,

 $Assistant\ Secretary\ for\ Aging. \\ [FR\ Doc.\ 2010-11202\ Filed\ 5-11-10;\ 8:45\ am]$ 

BILLING CODE 4154-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Health Resources and Services Administration

Health Care Integrity and Protection Data Bank (HIPDB) and National Practitioner Data Bank (NPDB): Public Posting of Non-Compliant Government Agencies

**AGENCY:** Health Resources and Services Administration, HHS.

**ACTION:** Notice of intent to publish list of non-compliant Government agencies.

SUMMARY: "Government agencies," as defined in section 1128E(g)(3) of the Social Security Act (42 U.S.C. 1320a–7e(g)(3)), that are not in compliance with the reporting requirements of the HIPDB will have their names published in a report on the HRSA and Data Bank Web sites (http://www.hrsa.gov and http://www.npdb-hipdb.hrsa.gov) by July 1, 2010. This listing of noncompliant Government agencies will be reviewed and updated on a periodic basis.

SUPPLEMENTARY INFORMATION: The HIPDB was mandated by Section 1128E of the Social Security Act (SSA) as added by Section 221(a) of the Health Insurance Portability and Accountability Act of 1996. Government agencies that license or certify health care practitioners, providers or suppliers, must report final adverse actions to the HIPDB generally within 30 days of the date the action becomes final. With the March 1, 2010, effective date of the final rule implementing Section 1921 of the SSA, many of the actions reported to the HIPDB also are

now posted and available for querying in the NPDB.

Section 1128E(b)(6)(B) of the SSA (42 U.S.C. 1320a–7e(b)(6)(B)) states that "[t]he Secretary shall provide for a publication of a public report that identifies those Government agencies that have failed to report information on adverse actions as required to be reported [to the HIPDB]."

Beginning no later than July 1, 2010, the Department will publish a report on the HRSA and Data Bank Web sites that identifies those professions for which Government agencies' reporting history has been analyzed. The report will also set forth a list of those Government agencies that are: (1) Out of compliance with reporting requirements (that is, they have failed to address their noncompliance); and (2) working toward full compliance with reporting requirements (that is, they have begun reporting). The listing of non-compliant Government agencies will be reviewed and updated on a periodic basis.

## FOR FURTHER INFORMATION CONTACT:

Mark S. Pincus, Acting Director, Division of Practitioner Data Banks, Bureau of Health Professions, Health Resources and Services Administration, 5600 Fishers Lane, Suite 8–103, Rockville, Maryland 20857. Tel: (301) 443–2300.

Dated: May 5, 2010.

## Mary K. Wakefield,

Administrator.

[FR Doc. 2010-11368 Filed 5-11-10; 8:45 am]

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# 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: National Institute of Allergy and Infectious Diseases Special

Emphasis Panel; TB Immunology and Drug Discovery.

Date: June 2, 2010.

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

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6700B Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Annie Walker-Abbey, PhD, Scientific Review Officer, Scientific Review Program, NIAID/NIH/DHHS, 6700B Rockledge Drive, RM 3126, MSC-7616, Bethesda, MD 20892-7616, 301-451-2671, aabbey@niaid.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 6, 2010.

# Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–11359 Filed 5–11–10; 8:45 am]

BILLING CODE 4140-01-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: Microbiology, Infectious Diseases and AIDS Initial Review Group; Microbiology and Infectious Diseases Research Committee.

Date: June 17, 2010.

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

Agenda: To review and evaluate grant applications.

*Place:* The Allerton Hotel, 701 North Michigan Avenue, Chicago, IL 60611.

Contact Person: Michelle M. Timmerman, PhD, Scientific Review Officer, Scientific Review Program, DEA/NIAID/NIH/DHHS, Room 2217, 6700B Rockledge Drive, MSC-7616, Bethesda, MD 20892–7616, 301–451–4573, timmermanm@niaid.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 6, 2010.

#### Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010-11323 Filed 5-11-10; 8:45 am]

BILLING CODE 4140-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### **National Institutes of Health**

### National Center for Research Resources; 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 Center for Research Resources Initial Review Group; Comparative Medicine Review Committee CMRC 2.

Date: June 2, 2010.

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

Agenda: To review and evaluate grant applications.

*Place:* Hilton Washington/Rockville, 1750 Rockville Pike, Rockville, MD 20852.

Contact Person: Bonnie B. Dunn, PhD, Scientific Review Officer, National Center for Research Resources, or National Institutes of Health, 6701 Democracy Blvd., 1 Democracy Plaza, Room 1074, MSC 4874, Bethesda, MD 20892–4874, 301–435–0824, dunnbo@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research; 93.371, Biomedical Technology; 93.389, Research Infrastructure, 93.306, 93.333; 93.702, ARRA Related Construction Awards, National Institutes of Health, HHS)

Dated: May 6, 2010.

## Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–11319 Filed 5–11–10; 8:45 am]

BILLING CODE 4140-01-P