topic. At least 7 minutes will be allotted for each speaker, and if time permits, may be extended up to 10 minutes at the discretion of the chair. Registration for oral comments will also be available on-site, although time allowed for presentation by on-site registrants may be less than for registered speakers and will be determined by the number of persons who register at the meeting. In addition to in-person oral comments at the meeting, public comments can be presented by teleconference line. There will be 50 lines for this call; availability will be on a first-come, first-served basis. The lines will be open from 8:30 a.m. until approximately 5:00 p.m., although public comments will be received only during the formal public comment periods, which will be indicated on the preliminary agenda. The access number for the teleconference line will be provided to registrants by email prior to the meeting.

Persons wishing to present oral comments are encouraged to register using the SACATM meeting registration form (*http://ntp.niehs.nih.gov/go/32822*), indicate the topic(s) on which they plan to comment, and, if possible, send a copy of their statement to *whiteld@niehs.nih.gov* by August 26, 2015, to enable review by SACATM, NICEATM, ICCVAM, and NIEHS/NTP staff prior to the meeting. Written statements can supplement and may expand the oral presentation. If registering on-site and reading from written text, please bring 20 copies of the statement for distribution and to supplement the record.

Background Information on ICCVAM, NICEATM, and SACATM: ICCVAM is an interagency committee composed of representatives from 15 federal regulatory and research agencies that require, use, generate, or disseminate toxicological and safety testing information. ICCVAM conducts technical evaluations of new, revised, and alternative safety testing methods with regulatory applicability and promotes the scientific validation and regulatory acceptance of toxicological and safety-testing methods that more accurately assess the safety and hazards of chemicals and products and that reduce, refine (decrease or eliminate pain and distress), or replace animal use. The ICCVAM Authorization Act of 2000 (42 U.S.C. 285*l*–3) established ICCVAM as a permanent interagency committee of the NIEHS under NICEATM.

NICEATM administers ICCVAM, provides scientific and operational support for ICCVAM-related activities, and conducts independent validation studies to assess the usefulness and limitations of new, revised, and alternative test methods and strategies. NICEATM and ICCVAM work collaboratively to evaluate new and improved test methods and strategies applicable to the needs of U.S. federal agencies. NICEATM and ICCVAM welcome the public nomination of new, revised, and alternative test methods and strategies for validation studies and technical evaluations. Additional information about ICCVAM and NICEATM can be found at *http://ntp.niehs.nih.gov/go/iccvam* and *http://ntp.niehs.nih.gov/go/niceatm.*

SACATM was established in response to the ICCVAM Authorization Act [Section 285*l*–3(d)] and is composed of scientists from the public and private sectors. SACATM advises ICCVAM, NICEATM, and the Director of the NIEHS and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM. SACATM 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 *http://ntp.niehs.nih.gov/go/167.*

Dated: July 8, 2015.

John R. Bucher,

Associate Director, National Toxicology Program.

[FR Doc. 2015–17165 Filed 7–13–15; 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 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 of Allergy and Infectious Diseases Special Emphasis Panel; NIAID Clinical Trial Implantation Grant (R01).

Date: August 17, 2015.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: 5061 Fishers Lane, Conference Room 3F100, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Paul Roberts, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G22, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9823, Rockville, MD 20852, 240–669–5053, *paul.roberts@nih.gov*.

Name of Committee: National Institute of Allergy and Infectious Diseases Special Emphasis Panel; NIAID Clinical Trial Implantation Grant (R01).

Date: August 18, 2015.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: 5601 Fishers Lane, Conference Room 3F100, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Paul Roberts, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G22, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9823, Rockville, MD 20852, 240–669–5053, *paul.roberts@nih.gov*.

Name of Committee: National Institute of Allergy and Infectious Diseases Special Emphasis Panel; NIAID Clinical Trial Implantation Cooperative Agreement (U01).

Date: August 19, 2015.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: 5601 Fishers Lane, Conference Room 3F100, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Paul Roberts, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G22, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9823, Rockville, MD 20852, 240–669–5053, *paul.roberts@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: July 8, 2015.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–17160 Filed 7–13–15; 8:45 am]

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