to provide additional information to HRSA by October 24, 2017, 12:00 p.m. Eastern Time. Please see contact information below.

FOR FURTHER INFORMATION CONTACT: Anyone requesting information regarding the ACHDNC should contact Ann Ferrero, Maternal and Child Health Bureau (MCHB), HRSA, in one of three ways: (1) Send a request to the following address: Ann Ferrero, MCHB, HRSA 5600 Fishers Lane, Room 18N100C, Rockville, MD 20857; (2) call 301–443–3999; or (3) send an email to: *AFerrero@hrsa.gov.*

SUPPLEMENTARY INFORMATION: The ACHDNC provides advice to the Secretary of HHS on the development of newborn screening activities, technologies, policies, guidelines, and programs for effectively reducing morbidity and mortality in newborns and children having, or at risk for, heritable disorders. In addition, ACHDNC's recommendations regarding inclusion of additional conditions and inherited disorders for screening which have been adopted by the Secretary are then included in the Recommended Uniform Screening Panel (RUSP). Conditions listed on the RUSP constitute part of the comprehensive preventive health guidelines supported by HRSA for infants and children under section 2713 of the Public Health Service Act, codified at 42 U.S.C. 300gg–13. Under this provision, non-grandfathered health plans are required to cover screenings included in the HRSA-supported comprehensive guidelines without charging a co-payment, co-insurance, or deductible for plan years (*i.e.,* policy years) beginning on or after the date that is one year from the Secretary's adoption of the condition for screening.

The meeting agenda will include: (1) An update on states' progress toward the newborn screening timeliness goals outlined by the Committee; (2) a presentation on phase 2 of the spinal muscular atrophy evidence review; (3) presentations on newborn screening topics such as the clinical and public health impact of Severe Combined Immunodeficiency (SCID), carrier status in the context of newborn screening, and a review of long term follow up in newborn screening; and (4) updates from the Laboratory Standards and Procedures workgroup, Follow-up and Treatment workgroup, and Education and Training workgroup. The Committee will not be voting on a proposed addition of a condition to the RUSP. Agenda items are subject to change. The final meeting agenda will be available 2 days prior to the meeting on the Committee's Web site: *http://www.hrsa.gov/advisorycommittees/mchbadvisory/heritabledisorders.*

Members of the public will have the opportunity to provide comments. All comments are part of the official Committee record. To submit written comments or request time for an oral comment at the meeting, please register online by 11:59 p.m. Eastern Time on November 2, 2017, at *http://www.achdncmeetings.org/.* To ensure all individuals who have registered and requested time for oral comments are accommodated, the allocated time for comments may be limited. Individuals associated with groups or who plan to provide comments on similar topics may be asked to combine their comments and present them through a single representative. No audiovisual presentations are permitted. Written comments should identify the individual's name, address, email, telephone number, professional or organization affiliation, background or area of expertise (*i.e.,* parent, family member, researcher, clinician, public health, etc.) and the topic/subject matter.

Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify Ann Ferrero using the address and phone number above at least 10 days prior to the meeting.

Amy McNulty,

Acting Director, Division of the Executive Secretariat.

[FR Doc. 2017–22313 Filed 10–13–17; 8:45 am]

BILLING CODE 4165–15–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, 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; NIAID Clinical Trial Planning Grant (R34) and Implementation Cooperative Agreement (U01).

Date: November 6–9, 2017.

Time: 9:00 a.m. to 5:30 p.m.

Agenda: To review and evaluate grant applications.

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

Contact Person: Geetanjali Bansal, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G49, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9834, Bethesda, MD 20892–9834, (240) 669–5073, *geetanjali.bansal@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: October 10, 2017.

Natasha M. Copeland,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–22259 Filed 10–13–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, 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 Cancer Institute Special Emphasis Panel; Translational Studies on Adducts For Cancer Risk Identification and Prevention.

Date: November 8, 2017.

Time: 1:00 p.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Cancer Institute, Shady Grove, 9609 Medical Center Drive, Room 7W108, Rockville, MD 20850, (Telephone Conference Call).