

Agenda: To review and evaluate grant applications.

Place: Crowne Plaza Washington National Airport, 1489 Jefferson Davis Hwy, Arlington, VA 22202.

Contact Person: Ryan G Morris, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4205, MSC 7814, Bethesda, MD 20892, 301-435-1501, morrisr@csr.nih.gov.
(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393-93.396, 93.837-93.844, 93.846-93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: September 14, 2015.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015-23387 Filed 9-17-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of a meeting of the Muscular Dystrophy Coordinating Committee (MDCC).

The meeting will be open to the public and accessible by live webcast.

Name of Committee: Muscular Dystrophy Coordinating Committee.

Type of meeting: Open Meeting.

Date: November 13, 2015.

Time: 8:30 a.m. to 4:30 p.m. *Eastern Time*—Approximate end time.

Agenda: The purpose of this meeting is to bring together committee members, representing government agencies, patient advocacy groups, other voluntary health organizations, and patients and their families to update one another on progress relevant to the Action Plan for the Muscular Dystrophies and to coordinate activities and discuss gaps and opportunities leading to better understanding of the muscular dystrophies, advances in treatments, and reduced disease burden. Prior to the meeting, an agenda will be posted to the MDCC meeting registration Web site: <https://meetings.ninds.nih.gov/meetings/MDCC13Nov2015/>.

Registration: To register, please go to: <https://meetings.ninds.nih.gov/meetings/MDCC13Nov2015/>.

Webcast Live: For those not able to attend in person, this meeting will be webcast at: <http://videocast.nih.gov/>.

Place: Neuroscience Center, Conference Room C/D, 6001 Executive Boulevard, Rockville, Maryland 20852.

Contact Person: Glen H. Nuckolls, Ph.D., Executive Secretary, Muscular Dystrophy

Coordinating Committee, National Institute of Neurological Disorders and Stroke, NIH, 6001 Executive Boulevard, NSC 2203, Rockville, MD 20852, (301) 496-5739, glen.nuckolls@ninds.nih.gov.

Any member of the public interested in presenting oral comments to the committee may notify the Contact Person listed on this notice at least 10 days in advance of the meeting. Interested individuals and representatives of organizations may submit a letter of intent, a brief description of the organization represented, and a short description of the oral presentation. Only one representative of an organization may be allowed to present oral comments and if accepted by the committee, presentations may be limited to five minutes. Both printed and electronic copies are requested for the record. In addition, any interested person may file written comments with the committee by forwarding their statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

Attendance is limited to seating space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should inform the Contact Person listed above in advance of the meeting. All visitors must go through a security check at the meeting site to receive a visitor's badge. A valid, government issued photo ID must be presented before a visitor's badge can be issued. Further information can be found at the registration Web site: <https://meetings.ninds.nih.gov/meetings/MDCC13Nov2015/>.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS)

Dated: September 14, 2015.

Carolyn Baum,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015-23443 Filed 9-17-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

In Vitro to In Vivo Extrapolation for High Throughput Prioritization and Decision Making; Notice of Webinars and Public Workshop; Registration Information

SUMMARY: The National Toxicology Program (NTP) Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM) and the U.S. Environmental Protection Agency (EPA) announce the workshop, “In Vitro to In Vivo Extrapolation for

High Throughput Prioritization and Decision Making.” Attendees at the in-person workshop and four webinar presentations leading up to the workshop will discuss the state of the science and best practices for using *in vitro* to *in vivo* extrapolation (IVIVE) in a tiered risk decision context.

DATES: *Webinars:* October 7, 2015, at 11:00 a.m. Eastern Daylight Time (EDT); and November 4, 2015; December 2, 2015; and January 6, 2016; at 11:00 a.m. Eastern Standard Time (EST).

Webinar Registration: Deadline is two business days prior to each webinar.

Workshop: February 17-18, 2016, from 9:00 a.m. to approximately 5:00 p.m. (EST).

Workshop Registration: Deadline is February 5, 2016 at 5:00 p.m. (EST).

ADDRESSES:

Workshop Location: U.S. Environmental Protection Agency, 109 T.W. Alexander Dr., Durham, NC, 27709.

Web page: The preliminary agenda, registration, and other meeting materials will be available at <http://ntp.niehs.nih.gov/go/ivive-wksp-2016>.

FOR FURTHER INFORMATION CONTACT: Dr. Warren S. Casey, Director, NICEATM; email: warren.casey@nih.gov; telephone: (919) 316-4729.

SUPPLEMENTARY INFORMATION:

Background: Data from high throughput *in vitro* tests are being generated for many chemicals of environmental and commercial interest, with the expectation that *in vitro* assay data could ultimately be used to predict adverse effects of chemical exposures *in vivo*. Translating values obtained from *in vitro* assays into estimates of *in vivo* outcomes is a complex process involving the use of mathematical modeling and increasingly complex test systems. The series of four webinars and in-person workshop aim to address the capabilities and limitations of IVIVE within the context of risk decision-making.

The webinar series will present the current science, and the in-person workshop will facilitate discussions that follow-up and build on information presented in the webinars. During the workshop, participants will (1) review the state of the science to form recommendations on best practices for using IVIVE in chemical screening and risk-based decision making, (2) identify areas that require additional data and/or research, and (3) highlight examples of how best to apply IVIVE in a tiered risk decision-making strategy.

Preliminary Agenda and Other Meeting Information: A preliminary agenda and additional information will

be available at <http://ntp.niehs.nih.gov/go/ivive-wksp-2016>.

Meeting and Registration: This workshop is open to the public, free of charge, with attendance limited only by the space available. Registration is required to attend both the webinars and the workshop. Those persons attending the workshop should plan to participate in all four webinars. However, viewing the webinars does not require attendance at the workshop. Individuals who plan to attend the workshop must register at <http://ntp.niehs.nih.gov/go/ivive-wksp-2016> by February 5, 2016. Individuals who plan to participate in the webinars must register at <http://ntp.niehs.nih.gov/go/ivive-wksp-2016> two business days prior to the webinar date to ensure access. Please visit this Web page for the most current information about the webinars and workshop. For those who register, information about how to access the webinar will be emailed within two business days of each webinar.

Individuals with disabilities who need accommodation to participate in these events should contact Dr. Elizabeth Maull at phone: (919) 316-4668 or email: maull@niehs.nih.gov. TTY users should contact the Federal TTY Relay Service at (800) 877-8339. Requests should be made at least five business days in advance of the event. Visitor and security information for those attending the workshop can be found at <http://www2.epa.gov/aboutepa/about-epas-campus-research-triangle-park-rtp-north-carolina>.

Background Information on NICEATM: NICEATM conducts data analyses, workshops, independent validation studies, and other activities to assess new, revised, and alternative test methods and strategies. NICEATM also provides support for the Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM). The ICCVAM Authorization Act of 2000 (42 U.S.C. 2851-3) provides authority for ICCVAM and NICEATM in the development of alternative test methods. Information about NICEATM and ICCVAM is found at <http://ntp.niehs.nih.gov/go/niceatm> and <http://ntp.niehs.nih.gov/go/iccvam>, respectively.

Dated: September 14, 2015.

John R. Bucher,

Associate Director, National Toxicology Program.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; 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 Aging Special Emphasis Panel; Aging of the Lung.

Date: October 20, 2015.

Time: 3:00 p.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

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

Contact Person: Maurizio Grimaldi, MD, Ph.D., Scientific Review Officer, National Institute on Aging, National Institutes of Health, 7201 Wisconsin Avenue, Room 2c218, Bethesda, MD 20892, 301-496-9374, grimaldim2@mail.nih.gov.

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

Dated: September 14, 2015.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; United States and Global Human Influenza Surveillance in At-Risk Settings (NIAID)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health, has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information

collection was previously published in the **Federal Register** on April 9, 2015, page 19090 and allowed 60-days for public comment. One comment was received. However, it was not applicable to this data collection. The purpose of this notice is to allow an additional 30 days for public comment. The National Institute of Allergy and Infectious Diseases (NIAID), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA submission@omb.eop.gov* or by fax to 202-395-6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, or request more information on the proposed project, contact: Dr. Diane Post, Program Officer, Respiratory Diseases Branch, NIAID, NIH, 5601 Fishers Lane, Bethesda, MD or call non-toll-free number at 240-627-3348 or email your request, including your address to: postd@niaid.nih.gov. Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: United States and Global Human Influenza Surveillance in at-Risk Settings, 0925-NEW, National Institute of Allergies and Infectious Diseases (NIAID), National Institutes of Health (NIH).

Need and Use of Information Collection: These studies will identify individuals with or at risk for influenza through focused surveillance in at-risk settings within the United States and internationally, rapidly identify circulating influenza strains to identify those with pandemic potential and create an invaluable bank of human samples from influenza patients to allow the characterization of the determinants of influenza transmission to and among humans, the immune response to influenza, and the basis of severe disease—critical knowledge gaps impacting effectiveness of decision-making around patient care and