

Research Training and Medical Education, NIH Clinical Center, National Institutes of Health, 10 Center Drive, Room 1N252C, Bethesda, MD 20892-1158, or call non-toll-free number (301) 496-2636, or Email your request, including your address to: robert.lembo@nih.gov. Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency'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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Proposed Collection Title: Application Process for Clinical Research Training and Medical Education at the NIH Clinical Center, Revision OMB #0925-0698, Expiration date July 31, 2020, National Institutes of Health Clinical Center (CC), National Institutes of Health (NIH).

Need and Use of Information Collection: The primary objective of the application process is to allow the Office of Clinical Research Training and Medical Education (OCRTME) at the NIH Clinical Center to evaluate

applicants' qualifications to determine applicants' eligibility for training programs managed by the Office. Applicants must provide the required information requested in the respective applications to be considered a candidate for participation. Information submitted by candidates for training programs is reviewed initially by OCRTME administrative staff to establish eligibility for participation. Eligible candidates are then referred to the designated training program director/administrator or training program selection committee for review and decisions regarding acceptance for participation. A secondary objective of the application process is to track enrollment in training programs over time.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours 333.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Clinical Electives Program	Pre Doctoral Students	300	1	20/60	100
Graduate Medical Education	Physicians	100	1	20/60	33
Medical Research Scholars Program	Pre Doctoral Students	200	1	20/60	67
Resident Electives Program	Physicians	100	1	20/60	33
Bioethics Fellowship Program	Pre Doctoral, Post-Doctoral	300	1	20/60	100
Total	1,000	333

Dated: April 10, 2020.
Laura M. Lee,
Project Clearance Liaison, NIH Clinical Center, National Institutes of Health.
 [FR Doc. 2020-08016 Filed 4-15-20; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Interagency Coordinating Committee on the Validation of Alternative Methods; Notice of Public Webcast; Request for Public Input

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM) will hold a public forum to share information and facilitate direct communication of ideas and suggestions from stakeholders. Interested persons

may view the presentations by webcast. Time will be set aside for questions and public statements on the topics discussed. Registration is required for both webcast viewing and oral statements. Information about the meeting and registration are available at <https://ntp.niehs.nih.gov/go/iccvamforum-2020>.

DATES:

Webcast: May 21, 2020, 9:00 a.m. to approximately 3:00 p.m. EDT.

Registration for Webcast: April 17, 2020, until 3:00 p.m. EDT May 21, 2020.

Registration for Oral Statements: April 17, 2020, until 4:00 p.m. EDT May 8, 2020.

Registration to view the webcast and present oral public statements is required.

ADDRESSES:

Meeting web page: A preliminary agenda will be posted by May 1 at <https://ntp.niehs.nih.gov/go/iccvamforum-2020>.

Webcast: The meeting will be webcast; information to connect to the

webcast will be provided to those who register for viewing.

FOR FURTHER INFORMATION CONTACT: Dr. Nicole Kleinstreuer, Acting Director, National Toxicology Program Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM), Division of NTP, NIEHS, P.O. Box 12233, K2-17, Research Triangle Park, NC 27709. Phone: 984-287-3150, Email: nicole.kleinstreuer@nih.gov. Hand Deliver/Courier address: 530 Davis Drive, Room K2021, Morrisville, NC 27560.

SUPPLEMENTARY INFORMATION:

Background: ICCVAM, a congressionally mandated committee, promotes the development and validation of alternative testing strategies that protect human health and the environment while replacing, reducing, or refining animal use.

ICCVAM's goals include promotion of national and international partnerships between governmental and nongovernmental groups, including academia, industry, advocacy groups, and other key stakeholders. To foster

these partnerships ICCVAM initiated annual public forums in 2014 to share information and facilitate direct communication of ideas and suggestions from stakeholders (79 FR 25136).

This year's meeting will be held on May 21, 2020. Due to restrictions on in-person gatherings amid ongoing public health concerns, the public forum will be presented via webcast only. NICEATM and ICCVAM members will give presentations on current activities related to the development and validation of alternative test methods and approaches, including activities relevant to implementation of the strategic roadmap for establishing new approaches to evaluate the safety of chemicals and medical products in the United States (83 FR 7487).

There will be opportunities for registered participants to ask clarifying or follow-up questions of the ICCVAM members about their presentations. Instructions for submitting these questions will be provided via email prior to the webcast. The agenda will also include time for public oral statements relevant to the ICCVAM mission and current activities from participants who have registered to do so in advance.

Preliminary Agenda and Other Meeting Information: A preliminary agenda will be posted by May 1 at <https://ntp.niehs.nih.gov/go/iccvamforum-2020>. Interested individuals are encouraged to visit this web page to stay abreast of the most current meeting information.

Webcast and Registration: This webcast is open to the public. Registration for the webcast is required and is open through 3:00 p.m. EDT on May 21, 2020 at <https://ntp.niehs.nih.gov/go/commprac-2020>. Registrants will receive instructions on how to access and participate in the webcast in the email confirming their registration.

Individuals with disabilities who need accommodation to participate in this event should contact Elizabeth Maull at phone: (984) 287-3157 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.

Request for Oral Public Statements: In addition to time for clarifying or follow-up questions following scheduled presentations, time will be allotted during the meeting for oral public statements with associated slides on topics relevant to ICCVAM's mission. Separate registration for those wishing to provide public statements is required and is open through May 8, 2020 at

<https://ntp.niehs.nih.gov/go/commprac-2020>. Any participant registered for the webcast may ask clarifying questions during the appropriate times in the agenda. The additional registration is only required for those who wish to give separate public statements. The number and length of presentations may be limited based on available time. Submitters will be identified by their name and affiliation and/or sponsoring organization, if applicable. Persons submitting public statements and/or associated slides should include their name, affiliation (if any), mailing address, telephone, email, and sponsoring organization (if any) with the document. National Toxicology Program guidelines for public statements are at http://ntp.niehs.nih.gov/ntp/about_ntp/guidelines_public_comments_508.pdf.

Participants registered to present oral public statements must email their statement to ICCVAMquestions@niehs.nih.gov by May 8, 2020, to allow time for review by NICEATM and ICCVAM and posting to the meeting page prior to the forum. Persons presenting oral public statements will be contacted to arrange the logistics of their presentations. Written statements may supplement and expand the oral presentation. Public statements will be distributed to NICEATM and ICCVAM members before the meeting.

Responses to this notice are voluntary. No proprietary, classified, confidential, or sensitive information should be included in statements submitted in response to this notice or presented during the meeting. This request for input is for planning purposes only and is not a solicitation for applications or an obligation on the part of the U.S. Government to provide support for any ideas identified in response to the request. Please note that the U.S. Government will not pay for the preparation of any information submitted or for its use of that information.

Background Information on ICCVAM and NICEATM: ICCVAM is an interagency committee composed of representatives from 16 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 and integrated testing strategies with regulatory applicability and promotes the scientific validation and regulatory acceptance of testing methods that more accurately assess the safety and hazards of chemicals and products and replace, reduce, or refine (enhance animal well-

being and lessen or avoid pain and distress) animal use.

The ICCVAM Authorization Act of 2000 (42 U.S.C. 285l-3) establishes ICCVAM as a permanent interagency committee of NIEHS and provides the authority for ICCVAM involvement in activities relevant to the development of alternative test methods. Additional information about ICCVAM can be found at <https://ntp.niehs.nih.gov/go/iccvam>.

NICEATM administers ICCVAM, provides scientific and operational support for ICCVAM-related activities, and conducts and publishes analyses and evaluations of data from new, revised, and alternative testing approaches. NICEATM and ICCVAM work collaboratively to evaluate new and improved testing approaches 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 NICEATM can be found at <https://ntp.niehs.nih.gov/go/niceatm>.

Dated: April 7, 2020.

Brian R. Berridge,

Associate Director, National Toxicology Program.

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DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS-2020-0006]

Homeland Security Advisory Council; Meeting

AGENCY: The Office of Partnership and Engagement (OPE), The Department of Homeland Security (DHS).

ACTION: Notice of partially closed teleconference Federal Advisory Committee meeting.

SUMMARY: The Homeland Security Advisory Council ("HSAC" or "Council") will meet via conference call on Thursday, May 7, 2020. The meeting will be partially closed to the public. Members of the public will be in listen-only mode during the open session.

DATES: The Council conference call will take place from 9:15 a.m. to 12:45 p.m. EDT on Thursday, May 7, 2020. The meeting will be closed to the public from 9:15 a.m. to 11:35 a.m. EDT. The meeting will be open to the public from 11:40 a.m. to 12:45 a.m. EDT. Please note the meeting may end early if the Council has completed its business, and