Review, National Institutes of Health, 6701 Rockledge Drive, Room 4120, MSC 7802, Bethesda, MD 20892, (301) 806–7314, *shahb@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Topics in Virology.

Date: July 28, 2021.

Time: 10:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Marci Scidmore, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3192, MSC 7808, Bethesda, MD 20892, 301–435–1149, *marci.scidmore@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: HIV Molecular Virology, Cell Biology, Immune and Therapeutic-Focused Application.

Date: July 28, 2021.

Time: 10:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Audrey O. Lau, MPH, Ph.D., Chief Center for Scientific Review, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 594–4088, *audrey.lau@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Endocrinology, Metabolism, Nutrition and Reproductive Sciences.

Date: July 28, 2021.

Time: 10:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Elaine Sierra-Rivera, Ph.D., IRG Chief, EMNR IRG, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6182, MSC 7892, Bethesda, MD 20892, 301–435–2514, *riverase@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Social Sciences and Chronic and Infectious Disease Epidemiology.

Date: July 28, 2021.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Lisa Steele, Ph.D., Scientific Review Officer, PSE IRG, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3139, MSC 7770, Bethesda, MD 20892, 301–594–6594, *steeleln@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Social Sciences and Chronic and Infectious Disease Epidemiology.

Date: July 28, 2021.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Denise Wiesch, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3138, MSC 7770, Bethesda, MD 20892, (301) 437–3478, *wieschd@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; UNITE Transformative Research to Address Health Disparities and Advance Health Equity at Minority Serving Institutions (U01).

Date: July 29–30, 2021.

Time: 10:00 a.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Aruna K. Behera, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4211, MSC 7814, Bethesda, MD 20892, 301–435–6809, *beheraak@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Topics in Mycology, Parasitology and Pathogenesis.

Date: July 29, 2021.

Time: 10:00 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Kenneth M. Izumi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3204, MSC 7808, Bethesda, MD 20892, 301–496–6980, *izumikm@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Topics in Endocrinology and Metabolism.

Date: July 29, 2021.

Time: 10:30 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Liliana N. Berti-Mattera, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, RM 6158, MSC 7890, Bethesda, MD 20892, (301) 827–7609, *liliana.berti-mattera@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Project: Endocannabinoids as Therapeutic Targets.

Date: July 29, 2021.

Time: 1:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Peter B. Guthrie, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4142, MSC 7850, Bethesda, MD 20892, (301) 435–1239, *guthriep@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Innate and Adaptive Immunology.

Date: July 30, 2021.

Time: 10:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Kenneth A. Roebuck, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5214, MSC 7852, Bethesda, MD 20892, (301) 435–1166, *roebuckk@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Topics in Innate and Adaptive Immunology.

Date: July 30, 2021.

Time: 10:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Neerja Kaushik-Basu, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3198, MSC 7808, Bethesda, MD 20892, (301) 435–1742, *kaushikbasun@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: June 24, 2021.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–13942 Filed 6–29–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Scientific Advisory Committee on Alternative Toxicological Methods; Announcement of Meeting; Request for Comments

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: This notice announces the next meeting of the Scientific Advisory Committee on Alternative Toxicological Methods (SACATM). SACATM is a federally chartered external advisory group of scientists from the public and private sectors, including

representatives of regulated industry and national animal protection organizations. SACATM advises the Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM), the National Toxicology Program (NTP) Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM), and the Director of the National Institute of Environmental Health Sciences (NIEHS) and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM. This SACATM meeting will be a virtual meeting only and available to the public for remote viewing. Registration is required to access the virtual meeting and to present oral comments. Information about the meeting and registration are available at *https://ntp.niehs.nih.gov/go/32822.*

DATES:

Meeting: September 28, 2021, 10:00 a.m.–3:30 p.m. EDT; September 29, 2021, 10:00 a.m.–4:00 p.m. EDT. Ending times are approximate; meeting may end earlier or run later.

Registration for Virtual Meeting: Deadline is September 29, 2021, 4:00 p.m. EDT.

Written Public Comment Submissions: Deadline is September 17, 2021.

Register to Present Oral Comments: Deadline is September 17, 2021.

Registration to view the virtual meeting and present oral public comments is required.

ADDRESSES:

Meeting web page: The preliminary agenda, registration, and other meeting materials are at *https://ntp.niehs.nih.gov/go/32822.*

Virtual Meeting: The URL for viewing the virtual meeting will be provided to those who register for viewing.

FOR FURTHER INFORMATION CONTACT: Dr. Sheena Scruggs, Designated Federal Official for SACATM, Office of Policy, Review, and Outreach, Division of NTP, NIEHS, P.O. Box 12233, K2–03, Research Triangle Park, NC 27709. Phone: 984–287–3355, Email: *sheena.scruggs@nih.gov.* Hand Deliver/Courier address: 530 Davis Drive, Room K2126, Morrisville, NC 27560.

SUPPLEMENTARY INFORMATION:

Meeting and Registration: The meeting is open to the public with time scheduled for oral public comments. Due to restrictions on in-person gatherings amid ongoing public health concerns, the meeting will be convened as a virtual meeting.

SACATM will provide input to ICCVAM, NICEATM, and NIEHS on programmatic activities and issues. Preliminary agenda items for the upcoming meeting include: (1) Major ICCVAM accomplishments in 2021; (2) regulatory needs and research applications for ecotoxicity testing; (3) evolving approaches to validation; and (4) update on NICEATM computational resources. Please see the preliminary agenda for information about specific presentations.

The preliminary agenda, roster of SACATM members, background materials, public comments, and any additional information will be posted when available on the SACATM meeting website (*https://ntp.niehs.nih.gov/go/32822*) or may be requested in hardcopy from the Designated Federal Official for SACATM. Following the meeting, summary minutes will be prepared and made available on the SACATM meeting website.

Registration is required to attend the virtual meeting and is open to all interested persons. Registrants will receive instructions on how to access the virtual meeting in the email confirming their registration. Individuals who plan to provide oral comments (see below) are required to register online at the SACATM meeting website (*https://ntp.niehs.nih.gov/go/32822*) by September 17, 2021, to facilitate planning for the meeting. Individuals are encouraged to visit the website often to stay abreast of the most current information regarding the meeting.

Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify Ms. Robbin Guy at phone: (984) 287–3136 or email: *robbin.guy@nih.gov* in advance of the meeting. 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.

Written Public Comments: Written and oral public comments are invited for the agenda topics. Guidelines for public comments are available at *https://ntp.niehs.nih.gov/ntp/about_ntp/guidelines_public_comments_508.pdf.* The deadline for submission of written comments is September 17, 2021. Written public comments should be submitted through the meeting website. Persons submitting written comments should include name, affiliation, mailing address, phone, email, and sponsoring organization (if any). Written comments received in response to this notice will be posted on the NTP website, and the submitter will be identified by name, affiliation, and sponsoring organization (if any).

Oral Public Comment Registration: The preliminary agenda allows for several public comment periods, each allowing up to six commenters a maximum of five minutes per speaker. Registration for oral comments is on or before September 17, 2021, at *https://ntp.niehs.nih.gov/go/32822.* Registration is on a first-come, first-served basis. Each organization is allowed one time slot per comment period. After the maximum number of speakers per comment period is exceeded, individuals registering to submit an oral comment for the topic will be placed on a wait list and notified should an opening become available. Commenters will be notified after September 17, 2021, to provide logistical information for their presentations. If possible, oral public commenters should send a copy of their slides and/or statement or talking points to Robbin Guy by email: *guyr2@niehs.nih.gov* by September 17, 2021.

Meeting Materials: The preliminary meeting agenda will be posted when available on the meeting web page at *https://ntp.niehs.nih.gov/go/32822* and will be updated one week before the meeting. Individuals are encouraged to visit this web page often to stay abreast of the most current information regarding 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, NICEATM, and SACATM: ICCVAM is an interagency committee composed of representatives from 17 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. 285*l*–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 *http://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 *http://ntp.niehs.nih.gov/go/niceatm.*

SACATM, established by the ICCVAM Authorization Act [Section 285l–3(d)], 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 to ICCVAM, NICEATM, and Director of NIEHS and NTP. 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: June 24, 2021.

Brian R. Berridge,

Associate Director, National Toxicology Program.

[FR Doc. 2021–13919 Filed 6–29–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer at (240) 276–0361.

Project: 2022 National Survey on Drug Use and Health (OMB No. 0930–0110)

SAMHSA is requesting from the Office of Management and Budget (OMB) approval to administer the National Survey on Drug Use and Health (NSDUH), a survey of the U.S. civilian, non-institutionalized population aged 12 years old or older. NSDUH data are used to determine the prevalence of use of tobacco products, alcohol, illicit substances, and illicit use of prescription drugs. The results are used by SAMHSA, the Office of National Drug Control Policy (ONDCP), federal government agencies, and other organizations and researchers to establish policy, direct program activities, and better allocate resources.

As certain parts of the United States reduce COVID–19 restrictions, NSDUH in-person data collection will proceed where possible. However, to ensure sufficient data are collected to produce nationally representative estimates for the 2022 survey, NSDUH will continue to employ a mix of in-person and web-based modes of administration to allow those respondents living in areas with COVID–19 restrictions the opportunity to participate. If the COVID–19 pandemic subsides to such levels to allow in-person data collection to resume nationwide, SAMHSA may reassess that multimode data collection model as part of the 2022 NSDUH.

In those areas where in-person data collection is permitted, NSDUH protocols, processes, and materials will continue to reflect the need to ensure the safety of respondents and field interviewers with respect to COVID–19—after initial implementation of such measures beginning in October 2020—which include equipping field interviewers with masks, gloves, disinfecting wipes, and hand sanitizer for use during data collection and providing a COVID–19 risk information form to all respondents.

Unlike previous NSDUHs, a hybrid address-based sampling (ABS) design will be implemented for the 2022 NSDUH. ABS refers to the sampling of residential addresses from a list based on the U.S. Postal Service's Computerized Delivery Sequence file. In areas with high expected ABS coverage, the ABS frame will be used. In all other areas, traditional field enumeration will be used to construct the dwelling unit frames.

In addition, the NSDUH questionnaire must be updated periodically to reflect changing substance use and mental health issues and to continue producing current data. For the 2022 NSDUH, the following questionnaire updates are planned: (1) Replacing the tobacco module with a redesigned nicotine module that includes questions about vaping, removes low priority items to reduce respondent burden and eliminates outdated terminology; (2) revising the marijuana module to include questions about the use of CBD, update questions on the mode of administration and eliminate outdated terminology and includes changes to the market information for marijuana questions; (3) redesigning the adult and youth mental health services utilization modules into one Mental Health Service Utilization model to remove questions with outdated terminology and include questions about newer treatments with recent increases in popularity; and (4) replacing the drug treatment module with a redesigned alcohol and drug treatment module that includes questions about newer treatments and those that have increased in popularity, as well as eliminating outdated terminology and reducing respondent burden.

As with all NSDUH/NHSDA [1] surveys conducted since 1999, the sample size of the NSDUH main study for 2022 will be sufficient to permit prevalence estimates for each of the fifty states and the District of Columbia. The total annual burden estimate for the NSDUH main study is shown below in Table 1.

[1] Prior to 2002, the NSDUH was referred to as the National Household Survey on Drug Abuse (NHSDA).