4A(iii) of DP5 OD026428–04, Slide 16 of NRI Seminar 2020, and Figure C(i) of the "Generation and Characterization of Ebf3Null Alleles" section of Poster 2020; specifically, Western blot data from three (3) mice were falsely represented as data from five (5) mice;

• the hindlimb splay measurements of  $Ebf3^{+/+}$  and  $Ebf3^{+/-}$  mice in RR6 2019 and Figure 4D(ii) of DP5 OD026428–04 by changing the severity score of the splay measurements in male and female mice to falsely show enhanced severity of the dystonia symptoms in  $Ebf3^{+/-}$  mice;

Respondent entered into a Voluntary Settlement Agreement (Agreement). Respondent neither admits nor denies ORI's findings of research misconduct. This settlement is not an admission of liability on the part of Respondent. Respondent voluntarily agreed to the following:

(1) Respondent will have his research supervised for a period of three (3) years beginning on May 14, 2024 (the "Supervision Period"). Prior to the submission of an application for PHS support for a research project on which Respondent's participation is proposed and prior to Respondent's participation in any capacity in PHS-supported research, Respondent will submit a plan for supervision of Respondent's duties to ORI for approval. The supervision plan must be designed to ensure the integrity of Respondent's research. Respondent will not participate in any PHS-supported research until such a supervision plan is approved by ORI. Respondent will comply with the agreed-upon supervision plan.

(2) The requirements for Respondent's supervision plan are as follows:

i. A committee of 2–3 senior faculty members at the institution who are familiar with Respondent's field of research, but not including Respondent's supervisor or collaborators, will provide oversight and guidance for a period of three (3) years from the effective date of the Agreement. The committee will review primary data from Respondent's laboratory on a quarterly basis and submit a report to ORI at six (6) month intervals setting forth the committee meeting dates and Respondent's compliance with appropriate research standards and confirming the integrity of Respondent's research.

ii. The committee will conduct an advance review of each application for PHS funds, or report, manuscript, or abstract involving PHS-supported research in which Respondent is involved. The review will include a discussion with Respondent of the primary data represented in those

documents and will include a certification to ORI that the data presented in the proposed application, report, manuscript, or abstract are supported by the research record.

(3) During the Supervision Period, Respondent will ensure that any institution employing him submits, in conjunction with each application for PHS funds, or report, manuscript, or abstract involving PHS-supported research in which Respondent is involved, a certification to ORI that the data provided by Respondent are based on actual experiments or are otherwise legitimately derived and that the data, procedures, and methodology are accurately reported and not plagiarized in the application, report, manuscript, or abstract.

(4) If no supervision plan is provided to ORI, Respondent will provide certification to ORI at the conclusion of the Supervision Period that his participation was not proposed on a research project for which an application for PHS support was submitted and that he has not participated in any capacity in PHS-supported research.

(5) During the Supervision Period, Respondent will exclude himself voluntarily from serving in any advisory or consultant capacity to PHS including, but not limited to, service on any PHS advisory committee, board, and/or peer review committee.

Dated: May 23, 2024.

### Sheila Garrity,

Director, Office of Research Integrity, Office of the Assistant Secretary for Health.

[FR Doc. 2024–11829 Filed 5–29–24; 8:45 am]

BILLING CODE 4150-31-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 1009 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; Stimulating Access to Research in Residency (StARR) (R38 Independent Clinical Trial Not Allowed).

Date: July 15, 2024.

Time: 10:00 a.m. to 2:00 p.m. Agenda: To review and evaluate grant applications.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3G22, Rockville, MD 20852 (Video Assisted Meeting).

Contact Person: Richard G. Kostriken, Ph.D., A.B., B.A., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3G22, Rockville, MD 20852, 240–669–2075, richard.kostriken@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: May 24, 2024.

### Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–11857 Filed 5–29–24; 8:45 am]

BILLING CODE 4140-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## **National Institutes of Health**

# National Center for Complementary & Integrative Health; Notice of Closed Meeting

Pursuant to section 1009 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 Center for Complementary and Integrative Health Special Emphasis Panel; Exploratory Clinical Trials of Mind and Body Interventions (MB).

Date: June 27–28, 2024.

Time: 2:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Center for Complementary and Integrative, Democracy II, 6707 Democracy Blvd., Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Mei Qin, MD, Ph.D., Scientific Review Officer, Office of Scientific Review, Division of Extramural Activities, NCCIH/NIH, 6707 Democracy Blvd., Suite 401, Bethesda, MD 20892, mei.qin@nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.213, Research and Training in Complementary and Alternative Medicine, National Institutes of Health, HHS)

Dated: May 23, 2024.

#### David W. Freeman,

Supervisory Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024-11854 Filed 5-29-24; 8:45 am]

BILLING CODE 4140-01-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 1009 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 contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, 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; Development of Medical Countermeasures for Biodefense and Emerging Infectious Diseases, Research Area 003—In Vitro Diagnostics (N01).

Date: June 25–July 2, 2024. Time: 10:00 a.m. to 5:00 p.m. Agenda: To review and evaluate contract proposals.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3F52A, Rockville, MD 20852 (Video Assisted Meeting).

Contact Person: Shilpakala Ketha, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3F52A, Rockville, MD 20852, (301) 761–6821, shilpa.ketha@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: May 24, 2024.

### Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024-11858 Filed 5-29-24; 8:45 am]

BILLING CODE 4140-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **National Institutes of Health**

# Office of the Secretary; Notice of Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the Interagency Autism Coordinating Committee.

This will be a hybrid meeting held inperson and virtually and will be open to the public as indicated below. Individuals who plan to attend inperson or view the virtual meeting and need special assistance or other accommodations should notify the Contact Person listed below at least seven (7) business days prior to the meeting. The meeting can be accessed from the NIH Videocast website (http://videocast.nih.gov/).

Name of Committee: Interagency Autism Coordinating Committee.

Date: July 10, 2024.

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

Agenda: To discuss committee business, updates, and issues related to autism research and services activities.

Place: National Institute of Mental Health (NIMH), Neuroscience Center (NSC), 6001 Executive Boulevard, First Floor Conference Room, Rockville, MD 20852 (In Person and Virtual).

Cost: The meeting is free and open to the public.

Registration: A registration web link will be posted on the IACC website (www.iacc.hhs.gov) prior to the meeting. Preregistration is recommended.

Deadlines: Public Comment Due Date: Monday, June 24, by 5:00 p.m. ET, Public Comment Guidelines, For public comment instructions, see below.

Contact Person: Ms. Rebecca Martin, Office of National Autism Coordination, National Institute of Mental Health, NIH, Phone: 301–435–0886, Email: IACCPublicInquiries@mail.nih.gov.

Public Comments: The IACC welcomes written and oral/virtual public comments from members of the autism community and asks the community to review and adhere to its Public Comment Guidelines. In the 2021–2023 IACC Strategic Plan, the IACC lists the "Spirit of Collaboration"

as one of its core values, stating that, "We will treat others with respect, listen with open minds to the diverse lived experiences of people on the autism spectrum and their families, consider multiple solutions, and foster discussions where participants can comfortably share different opinions." In keeping with this core value, the IACC and the NIMH Office of National Autism Coordination (ONAC) ask that members of the public who provide public comments or participate in meetings of the IACC also adhere to this core value.

A limited number of slots are available for individuals to provide a ~3-minute summary or excerpt of their written comment to the Committee during the meeting either in person or via videoconference. For those interested in that opportunity, please indicate "Interested in providing oral/virtual comment" in your written submission, along with your name, address, email, phone number, and professional/organizational affiliation so that ONAC staff can contact you if a slot is available.

For any given meeting, priority for comment slots will be given to individuals and organizations that have not previously provided comments in the current calendar year. This will help ensure that as many individuals and organizations as possible have an opportunity to share comments. Commenters going over their allotted 3-minute slot may be asked to conclude immediately in order to allow other comments and the rest of the meeting to proceed on schedule.

Public comment submissions received by 5:00 p.m. ET on Monday, June 24, 2024, will be provided to the Committee prior to the meeting for their consideration. Any written comments received after 5:00 p.m. ET, Monday, June 24, 2024, may be provided to the Committee either before or after the meeting, depending on the volume of comments received and the time required to process them in accordance with privacy regulations and other applicable Federal policies. The Committee is not able to respond individually to comments. All public comments become part of the public record. Attachments of copyrighted publications are not permitted, but web links or citations for any copyrighted works cited may be provided. For public comment guidelines, see: https:// iacc.hhs.gov/meetings/publiccomments/guidelines/.

Technical issues: If you experience any technical problems with the webcast, please email IACCPublicInquiries@mail.nih.gov.