technical evaluations. Additional information about NICEATM can be found at *https://ntp.niehs.nih.gov/go/niceatm.*

SACATM, established by the ICCVAM Authorization Act [Section 285*l*-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. SACATM is governed by the provisions of the Federal Advisory Committee Act, as amended (5 U.S.C. ch.10), which sets forth standards for the formation and use of advisory committees.

Additional information about SACATM, including link to the charter, roster, and records of past meetings, can be found at *http://ntp.niehs.nih.gov/go/167.*

Dated: July 30, 2024.

Richard P. Woychik,

Director, National Institute of Environmental Health Sciences and National Toxicology Program, National Institutes of Health.

[FR Doc. 2024–17099 Filed 8–1–24; 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 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 Implementation Cooperative Agreement (U01 Clinical Trial Required).

Date: August 29, 2024.

Time: 10:00 a.m. to 1:00 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 3E71, Rockville, MD 20892 (Video Assisted Meeting).

Contact Person: Samita S. Andreansky, 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 3E71, Rockville, MD 20852, 240–669–2915, *samita.andreansky@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: July 29, 2024.

Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–17048 Filed 8–1–24; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; 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: Center for Scientific Review Special Emphasis Panel; Biomedical Technology Optimization and Dissemination Center (BTOD).

Date: September 26, 2024.

Time: 10:00 a.m. to 8:30 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: Joseph Thomas Peterson, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4118, MSC 7814, Bethesda, MD 20892, 301–408–9694, *petersonjt@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: July 29, 2024.

Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–17049 Filed 8–1–24; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Collaboration Opportunity for Combination of Vaccine With Adoptive Cell Therapies Made at NCI for the Treatment of Solid Cancers

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The Surgery Branch (SB) at the National Cancer Institute (NCI), is seeking a partner in the private sector to provide Good Manufacturing Practice-grade vaccine directed against cancer neo-antigens with the goal of conducting a Phase-I human clinical trial for solid cancers.

FOR FURTHER INFORMATION CONTACT: Inquiries relating to this collaboration opportunity should be directed to: Aida Cremesti, Ph.D., Senior Technology Transfer Manager, NCI Technology Transfer Center, Telephone: (240) 276–5530; Email: *aida.cremesti@mail.nih.gov.* Inquiries related to licensing the related technology E–046–2022 should be directed to: Andrew Burke, Ph.D., Senior Technology Transfer Manager, NCI Technology Transfer Center, Telephone: (240)–276–5484; Email: *andy.burke@nih.gov.*

SUPPLEMENTARY INFORMATION:

Collaboration Opportunity Summary

The Surgery Branch (SB) at the National Cancer Institute (NCI), under the direction of Dr. Steven Rosenberg, is seeking a partner in the private sector to provide a GMP-grade vaccine directed against cancer neo-antigens, either private (patient-specific neo-antigens) or shared common tumor antigens (such as KRAS or P–53), with the goal of conducting a Phase-I human clinical trial for solid cancers. The trial would involve the combination of NCI-engineered cell therapies with a vaccine to be provided by the partner. The NCI SB has extensive expertise in the latest technology of tumor infiltrating lymphocyte (TIL) development, as well as T-Cell Receptor (TCR)-transduced Peripheral Blood Lymphocytes (PBL) development using NCI proprietary