

would benefit from harmonization, consistency, clarity, simplification, and/or coordination.

The SACHRP meeting will open to the public at 8:30 a.m., on Tuesday, October 16, 2018, followed by opening remarks from Dr. Jerry Menikoff, Director of OHRP and Dr. Stephen Rosenfeld, SACHRP Chair.

The SAS and SOH subcommittees will present their recommendations regarding the description of “key information,” as required by the revised Common Rule at § 46.116(a)(5)(i). This will be followed by a discussion of recommendations of the interpretation of the revised Common Rule’s exemptions § 46.104(d)(1) and (2) for HHS funded research. Lastly, the committee will continue its July discussions in the Office of Inspector General Report, July 7, 2017: “OHRP Generally Conducted Its Compliance Activities Independently, But Changes Would Strengthen Its Independence.”

The Wednesday, October 17, meeting will begin at 8:30 a.m. The SAS subcommittee will present and discuss recommendations on the interpretation of “reasonably available” at § 46.408(b), as well as discuss issues surrounding payment of subjects for participation in research. Modifications to the previous day’s work will be discussed and finalized. The meeting will adjourn at approximately 4:00 p.m., October 17, 2018.

Time for public comment sessions will be allotted both days. On-site registration is required for participation in the live public comment session. Note that public comment must be relevant to topics currently being addressed by the SACHRP. Individuals submitting written statements as public comment should email or fax their comments to SACHRP at SACHRP@hhs.gov at least five business days prior to the meeting.

Public attendance at the meeting is limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify one of the designated SACHRP points of contact at the address/phone number listed above at least one week prior to the meeting.

Dated: September 14, 2018.

Julia G. Gorey,

Executive Director, Secretary’s Advisory Committee on Human Research Protections.

[FR Doc. 2018–20676 Filed 9–21–18; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Arthritis and Musculoskeletal and Skin Diseases; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings 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 Arthritis and Musculoskeletal and Skin Diseases Special Emphasis Panel; T32 Institutional Training Grants.

Date: October 10, 2018.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Canopy by Hilton, 940 Rose Avenue, North Bethesda, MD 20852.

Contact Person: Nakia C. Brown, Ph.D., Scientific Review Officer, 6701 Democracy Blvd., RM 816, Bethesda, MD 20892, 301–827–4905, brownnac@mail.nih.gov.

Name of Committee: Arthritis and Musculoskeletal and Skin Diseases Initial Review Group; Arthritis and Musculoskeletal and Skin Diseases Clinical Trials Review Committee.

Date: October 16–17, 2018.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Hilton Garden Inn, 7301 Waverly Street, Bethesda, MD 20814.

Contact Person: Nakia C. Brown, Ph.D., NIAMS/Scientific Review Officer, 6701 Democracy Blvd., RM 816, Bethesda, MD 20892, 301–827–4905, brownnac@mail.nih.gov.

Name of Committee: National Institute of Arthritis and Musculoskeletal and Skin Diseases Special Emphasis Panel; AMSC Member Conflict.

Date: October 29, 2018.

Time: 10:00 a.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Democracy One, 6701 Democracy Blvd., Bethesda, MD 20892.

Contact Person: Yasuko Furumoto, Ph.D., NIAMS/Scientific Review Officer, 6701 Democracy Blvd., RM 816, Bethesda, MD 20892, 301–451–6520, yasuko.furumoto@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.846, Arthritis,

Musculoskeletal and Skin Diseases Research, National Institutes of Health, HHS)

Dated: September 18, 2018.

Sylvia L. Neal,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2018–20666 Filed 9–21–18; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The invention listed below is owned by an agency of the U.S. Government and is available for licensing to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

FOR FURTHER INFORMATION CONTACT: Yogikala Prabhu, Ph.D., 301–761–7789; prabhuyo@niaid.nih.gov. Licensing information and copies of the patent applications listed below may be obtained by communicating with the indicated licensing contact at the Technology Transfer and Intellectual Property Office, National Institute of Allergy and Infectious Diseases, 5601 Fishers Lane, Rockville, MD 20852; tel. 301–496–2644. A signed Confidential Disclosure Agreement will be required to receive copies of unpublished patent applications.

SUPPLEMENTARY INFORMATION: Technology description follows.

A New Class of Immunomodulatory Drugs for Multiple Sclerosis

Description of Technology: Multiple sclerosis (MS) is an autoimmune disease caused by activated autoimmune T lymphocytes in patients resulting in inflammatory demyelination in the central nervous system. Current treatments are focused on functional control of these activated autoimmune T cells, but these treatments are non-specific T cell inhibitors and have serious, sometimes fatal side effects. A specific therapy aimed at eliminating these autoimmune T cells through restimulation-induced cell death (RICD) could cure the disease and overcome the fatal side effects of current therapies.

NIAID inventors have identified a multi-valent tolerogen (MMPt), which can specifically elicit RICD of the activated, disease causing autoimmune T cells without compromising the general T cell-dependent immunity in the host. Animal studies have demonstrated that MMPt exerts robust therapeutic effects on both monophasic as well as relapsing-remitting type of the disease, indicating its medical applicability for treating MS patients with active disease. NIAID is seeking partners to develop this multi-valent peptide to improve its efficacy for use in clinical trials.

This technology is available for licensing for commercial development in accordance with 35 U.S.C. 209 and 37 CFR part 404, as well as for further development and evaluation under a research collaboration.

Potential Commercial Applications:

- Therapeutics

Competitive Advantages:

- Tolerogen induced elimination of activated autoimmune T cells will overcome the fatal side effects of current therapies
- Treatment of all types of MS patients

Development Stage:

- Preclinical (In vitro and in vivo animal studies)

Inventors: Dr. Michael J. Lenardo (NIAID), Dr. Lixin Zheng (NIAID), Dr. Jian Li (NIAID), Dr. Jae Lee (NIAID), and Dr. Wei Lu (NIAID).

Intellectual Property: HHS Reference No. E-064-2015, U.S. Provisional Patent Application Numbers: 62/130,285 filed March 9, 2015 and 62/219,851 filed September 17, 2015, and US PCT application PCT/US2016/021571 filed on March 9, 2016. Entered National Stage filing in US, EU, Canada, and Australia.

Licensing Contact: Yogikala Prabhu, Ph.D., 301-761-7789; prabhuyo@niaid.nih.gov.

Collaborative Research Opportunity: The National Institute of Allergy and Infectious Diseases is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate or commercialize the MMPt peptide to improve its efficacy for use in clinical trials.

For collaboration opportunities, please contact Yogikala Prabhu, Ph.D., 301-761-7789; prabhuyo@niaid.nih.gov.

Dated: September 18, 2018.

Suzanne M. Frisbie,

Deputy Director, Technology Transfer and Intellectual Property Office, National Institute of Allergy and Infectious Diseases.

[FR Doc. 2018-20664 Filed 9-21-18; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Advisory Committee on Research on Women's Health.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting. The meeting will also be videocast and can be accessed from the NIH Videocasting and Podcasting website (<http://videocast.nih.gov/>).

Name of Committee: Advisory Committee on Research on Women's Health.

Date: October 23, 2018.

Time: 9:00 a.m. to 4:00 p.m.

Agenda: Opening Remarks, Director's Report, Scientific Presentations, ORWH SCORE Program Update, and Strategic Plan Launch.

Place: National Institutes of Health, Building 31, 6th Floor, Conference Room 10, 31 Center Drive, Bethesda, MD 20892.

Contact Person: Elizabeth Spencer, R.N., Deputy Director, Office of Research on Women's Health, Executive Secretary, ACRWH, National Institutes of Health, 6707 Democracy Blvd., Room 7W444, Bethesda, MD 20817, 301-402-1770, elizabeth.spencer@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.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

Information is also available on the Institute's/Center's home page: <https://orwh.od.nih.gov/>, where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.14, Intramural Research Training Award; 93.22, Clinical Research Loan Repayment Program for Individuals from Disadvantaged Backgrounds; 93.232, Loan Repayment Program for Research Generally; 93.39, Academic Research Enhancement Award; 93.936, NIH Acquired Immunodeficiency Syndrome Research Loan Repayment Program; 93.187, Undergraduate Scholarship Program for Individuals from Disadvantaged Backgrounds, National Institutes of Health, HHS)

Dated: September 18, 2018.

Ronald J. Livingston, Jr.,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2018-20665 Filed 9-21-18; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the NIH Clinical Center Research Hospital Board.

The meeting will be open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

Name of Committee: NIH Clinical Center Research Hospital Board.

Date: October 19, 2018.

Time: 9:00 a.m. to 3:25 p.m.

Agenda: Discussion of Patient Safety.

Place: National Institutes of Health, Building 31, 6th Floor, Room: 6C6, 31 Center Drive, Bethesda, MD 20892.

Contact Person: Gretchen Wood, Staff Assistant, Office of the Director, National Institutes of Health, One Center Drive, Building 1, Bethesda, MD 20892, 301-496-4272, woodgs@nih.gov.