DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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; Opportunities for Collaborative Research at the NIH Clinical Center (U01).

Date: July 1–2, 2013.

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

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge 6700, 6700B Rockledge Drive, Bethesda, MD 20817 (Telephone Conference Call).

Contact Person: B. Duane Price, Ph.D., Scientific Review Officer, Scientific Review Program, DHHS/NIH/NIAID, 6700B Rockledge Drive, MSC 7616, Room 3139, Bethesda, MD 20892, (301) 451–2592, *pricebd@niaid.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: June 4, 2013.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–13621 Filed 6–7–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Complementary and Alternative Medicine; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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 Alternative Medicine Special Emphasis Panel, Clinical Grants.

Date: July 9, 2013.

Time: 2:30 p.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892.

Contact Person: Hungyi Shau, Ph.D., Scientific Review Officer, National Center for Complementary and Alternative Medicine, National Institutes of Health, 6707 Democracy Boulevard, Suite 401, Bethesda, MD 20892, 301–402–1030, *Hungyi.Shau@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: June 4, 2013.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–13619 Filed 6–7–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Start-up Exclusive Evaluation License: Portable Device and Method for Detecting Hematomas

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: This is notice, in accordance with 35 U.S.C. 209(c)(1) and 37 CFR 404.7(a)(1)(i), that the National Institutes of Health (NIH), Department of Health and Human Services, is contemplating the grant of a worldwide exclusive evaluation option license to practice the inventions embodied in: HHS Ref. No. E–010–2010/0, U.S. Provisional Patent Application No. 61/286,626, filed December 15, 2009, International Patent Application PCT/US2010/060506 filed December 15, 2010 (published as WO2011084480), European Patent Application 10798422.1 filed December 15, 2010, and U.S. Patent Application 13/516,480 filed June 15, 2012, to ArcheOptix, having its principle place of business in Kingston, Ontario (Canada).

The United States of America is an assignee to the patent rights of these inventions.

The contemplated exclusive license may be limited to devices for the detection of hematomas. Upon the expiration or termination of the start-up exclusive evaluation license, ArcheOptix will have the right to execute a start-up exclusive patent commercialization license with no greater field of use and territory than granted in the evaluation license.

DATES: Only written comments and/or applications for a license that are received by the NIH Office of Technology Transfer on or before June 25, 2013 will be considered.

ADDRESSES: Requests for a copy of the patent application, inquiries, comments and other materials relating to the contemplated license should be directed to: Michael Shmilovich, Esq., Senior Licensing and Patent Manager, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852–3804; Telephone: (301) 435–5019; Facsimile: (301) 402–0220; Email: *shmilovm@mail.nih.gov*.

SUPPLEMENTARY INFORMATION: The invention is a device and method for detecting hematomas based on near infrared light emitted perpendicularly into a tissue from a non-stationary emitter and on continuous detection of the reflected light with a non-stationary probe. The device is designed as a handheld detector that can be used either in an ER or at the scene of an accident, which will allow the Doctor or EMT to diagnose hematoma for patients with a traumatic brain injury at the scene. Furthermore, this device can be utilized to discriminate between subdural, epidural and bi-lateral hematomas. The specific combination and sequences of data analysis are performed to discriminate healthy tissue from tissue perfused with blood. This invention will result in a better triage and treatment for patients with traumatic brain injury (TBI) and fills a must filled gap in TBI health care.

The prospective exclusive evaluation option license is being considered under the small business initiative launched on October 1, 2011 and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective exclusive evaluation option license, and a subsequent exclusive patent commercialization license, may