Contact Person: Reigh-Yi Lin, Ph.D., Scientific Review Officer, Center for Scientific Review, 6701 Rockledge Drive, Bethesda, MD 20892, 301–827–6009, lin.reigh-yi@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR 16–366—Dual Purpose with Dual Benefit: Research in Biomedicine and Agriculture.

Date: March 21, 2017.
Time: 8:00 a.m. to 6:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Tera Bounds, DVM, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3214, MSC 7808, Bethesda, MD 20892, 301–435–2306, boundst@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Risk, Prevention and Health Behavior Overflow.

Date: March 21, 2017.
Time: 9:00 a.m. to 1:00 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Stacey C. FitzSimmons, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3114, MSC 7808, Bethesda, MD 20892, (301) 451–9956, fitzsimmons@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Urology and Urogynecology Small Business Applications.

Date: March 21, 2017.
Time: 9:00 a.m. to 12:00 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Ganesan Ramesh, Ph.D., Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr., Room 2182, MSC 7818, Bethesda, MD 20892, ganesan.ramesh@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Topics in drug discovery and clinical field studies.

Date: March 21, 2017.
Time: 10:00 a.m. to 5:00 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Liangbiao Zheng, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3202, MSC 7808, Bethesda, MD 20892, 301–996–5819, zhenglb@csr.nih.gov.


Nathasha M. Copeland,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–03281 Filed 2–17–17; 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 inventions listed below are owned by an agency of the U.S. Government.

FOR FURTHER INFORMATION CONTACT: Licensing information may be obtained by emailing the indicated licensing contact at the National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 4A29, MSC 7808, Bethesda, MD 20892; telephone: 301–402–5579. A signed Confidential Disclosure Agreement may be required to receive any unpublished information.

SUPPLEMENTARY INFORMATION: The following inventions are available for licensing in accordance with 35 U.S.C. 209 and 37 CFR part 404 to achieve expeditious commercialization of results of federally-funded research and development. Technology description follows.

Methods for Improving Drug Delivery to the Central Nervous System

The invention relates to the uses of the tricyclic antidepressant amitriptyline, its bioactive metabolites, and other LPA1R activators to improve the bioavailability and delivery of therapeutics to the central nervous system. This invention demonstrates that amitriptyline and other agents selectively decrease P-glycoprotein (P-gp) transport activity by ligand activation of lysophosphatidic acid 1 receptor (LPA1R) at the blood-brain barrier. P-gp is an effective target for increasing drug delivery to the brain (CNS) for two major reasons: (1) Its substrates include a large portion of on-the-market drugs, including chemotherapeutics, and (2) its directionality results in a net efflux of drugs from the brain. Additionally, specifically targeting P-gp through LPA1R activation bypasses the clinical challenges resulting from the toxicity of substrate inhibitors of P-gp. This invention describes the inhibition of drug efflux by P-gp transport; thus, co-administration of therapeutics with amitriptyline and other LPA1R activators provides a method for increasing drug delivery to the CNS, and improving overall drug efficacy. Moreover, drug delivery to other barrier tissues will also be enhanced where a similar LPA1R–P-gp activity relationship exists.

Potential Commercial Applications:
• Drug Delivery to the CNS.
• Co-administration of therapeutics.
• Blood-brain-barrier permeability.

Development Stage:
• Early stage.

Inventors: Ronald Cannon and David Banks (NIH).

Publications:


Licensing Contact: Michael Shmiliovich, Esq., CLP; 301–435–5019; shmiliovich@mail.nih.gov.


Michael Shmiliovich,
Senior Licensing and Patenting Manager, National Heart, Lung, and Blood Institute, Office of Technology Transfer and Development.

[FR Doc. 2017–03306 Filed 2–17–17; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; Notice of Closed Meetings

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 meetings.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**