

2019, Bethesda, MD 20892, 301-443-2861, marmillotp@mail.nih.gov.

Name of Committee: National Institute on Alcohol Abuse and Alcoholism Initial Review Group; Neuroscience Review Subcommittee.

Date: October 27, 2016.

Time: 8:30 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute on Alcohol Abuse and Alcoholism, National Institutes of Health, Terrace Level Conference Room 508-509, 5635 Fishers Lane, Rockville, MD 20852.

Contact Person: Beata Buzas, Ph.D., Scientific Review Officer, National Institute On Alcohol Abuse and Alcoholism, National Institutes of Health, 5635 Fishers Lane, Rm. 2081, Rockville, MD 20852, 301-443-0800, bbuzas@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants; 93.701, ARRA Related Biomedical Research and Research Support Awards, National Institutes of Health, HHS)

Dated: August 2, 2016.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016-18688 Filed 8-5-16; 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, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: The invention listed below is owned by an agency of the U.S. Government and is available for licensing and/or co-development in the U.S. 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. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing and/or co-development.

ADDRESSES: Invention Development and Marketing Unit, Technology Transfer Center, National Cancer Institute, 9609 Medical Center Drive, Mail Stop 9702, Rockville, MD 20850-9702.

FOR FURTHER INFORMATION CONTACT: Information on licensing and co-

development research collaborations, and copies of the U.S. patent applications listed below may be obtained by contacting: Attn. Invention Development and Marketing Unit, Technology Transfer Center, National Cancer Institute, 9609 Medical Center Drive, Mail Stop 9702, Rockville, MD, 20850-9702, Tel. 240-276-5515 or email ncitechtransfer@mail.nih.gov. A signed Confidential Disclosure Agreement may be required to receive copies of the patent applications.

SUPPLEMENTARY INFORMATION:

Technology description follows.

Title of invention: Synthetic Human-Derived Peptides and Peptidomimetics for Cancer Therapeutics

Keywords: Rpn13, selective proteasome inhibitor, proteasome ubiquitin receptors, (competition: carfilzomib and bortezomib too toxic, resistance developed), solid tumors, hematological cancer, HPV associated cancer, ovarian cancer, prostate cancer, gastric cancer, breast cancer, or colorectal cancer

Description of Technology: FDA approved 26S proteasome inhibitors, such as carfilzomib and bortezomib (Velcade®) have proven to be effective at treating hematologic cancers. However, resistance to these agents as well as their toxicity have raised concerns and highlight the need for new 26S proteasome inhibitors.

Investigators at the NCI's Structural Biophysics Laboratory have developed a new class of proteasome inhibitors. They are hRpn2-derived peptides capable of specifically targeting the Pru domain of hRpn13. Disruption of the Rpn2/Rpn13 interaction inhibits proteolysis by a mechanism that differs from those of the approved proteasome inhibitors.

Potential Commercial Applications:

- New class of proteasome inhibitors, targeting hRpn13 of the regulatory particle.

Value Proposition:

- Synergistic with, and more specific than, known proteasome inhibitors.
- Alternate mechanism of action compared to approved proteasome inhibitors.

Development Stage: Discovery (Lead ID).

Inventor(s): Kylie J. Walters (NCI), Fen Liu (NCI), and Xiuxiu Lu (NCI).

Intellectual Property:

HHS Reference No. E-278-2015/0-US-01.

US Provisional Application 62/222,530 (HHS Reference No. E-278-2015) filed September 23, 2015 entitled "Human RPN2 Derived Peptides Useful For Treating Cancer".

Publications: 1. Lu X., et al., 2015 PLoS One 2015 Oct 14;10(10) PMID: 26466095.

Contact Information: Requests for copies of the patent application or inquiries about licensing, research collaborations, and co-development opportunities should be sent to John D. Hewes, Ph.D., email: john.hewes@nih.gov.

Dated: July 26, 2016.

John D. Hewes,

Technology Transfer Specialist, Technology Transfer Center, National Cancer Institute.

[FR Doc. 2016-18689 Filed 8-5-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App. 2), notice is hereby given of the meeting of the National Cancer Advisory Board (NCAB).

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 open session will be videocast and can be accessed from the NIH Videocasting and Podcasting Web site (<http://videocast.nih.gov>).

A portion of the National Cancer Advisory Board 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 Cancer Advisory Board.

Open: September 7, 2016, 9:00 a.m. to 05:00 p.m.

Agenda: Programs reports and presentations; business of the Board.

Place: National Institutes of Health, Building 31, C-Wing, 6th Floor, Room 10, 9000 Rockville Pike, Bethesda, MD 20892.

Closed: September 7, 2016, 04:00 p.m. to 05:00 p.m.

Agenda: Review of NCAB grant applications.

Place: National Institutes of Health, Building 31, C-Wing, 6th Floor, Room 10, 9000 Rockville Pike, Bethesda, MD 20892.

Contact Person: Paulette S. Gray, Ph.D., Director, Division of Extramural Activities, National Cancer Institute—Shady Grove, National Institutes of Health, 9609 Medical Center Drive, Room 7W444, Bethesda, MD 20892, 240-276-6340, grayp@mail.nih.gov.

Any interested person may file written comments with the committee by forwarding the 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, NCAB: <http://deainfo.nci.nih.gov/advisory/ncab/ncab.htm>, where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: August 2, 2016.

Melanie J. Gray,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016-18686 Filed 8-5-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; 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: Center for Scientific Review Special Emphasis Panel; Member Conflicts: Consequences of Aging.

Date: August 15, 2016.

Time: 3:30 p.m. to 6:00 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: Pat Manos, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5200, MSC 7846, Bethesda, MD 20892, 301-408-9866, manospa@csr.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(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: August 2, 2016.

Anna Snouffer,

Deputy Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2016-18687 Filed 8-5-16; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Modification of the National Customs Automation Program (NCAP) Test Concerning the Automated Commercial Environment (ACE) Portal Accounts To Establish the Protest Filer Account and Clarification That the Terms and Conditions for Account Access Apply to All ACE Portal Accounts

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: General notice.

SUMMARY: This document announces U.S. Customs and Border Protection's (CBP's) plan to modify the National Customs Automation Program (NCAP) test concerning Automated Commercial Environment (ACE) Portal Accounts to establish the ACE Protest Filer Account. After CBP deploys the ACE Protest Module test at a later date, participants with an ACE Protest Filer Account will be able to file an electronic protest in ACE. This document also clarifies that CBP's previously published terms and

conditions governing access to and use of the NCAP test of ACE Portal Accounts apply to all ACE Portal Accounts, including all ACE Portal Account types created after the previously published terms and conditions. All other aspects of the ACE Portal Accounts Test remain the same as set forth in previously published **Federal Register** notices.

DATES: The modifications and clarifications of the ACE Portal Account Test made by this notice are effective on August 8, 2016.

The clarification to the terms and conditions applies to all ACE Portal Accounts regardless of when the account was created.

ADDRESSES: Comments concerning this notice and any aspect of the modified ACE Portal Account Test may be submitted at any time during the testing period via email to Josephine Baiamonte, ACE Business Office (ABO), Office of Trade at josephine.baiamonte@cbp.dhs.gov. In the subject line of your email, please indicate, "Comment on ACE Portal Account Test FRN".

FOR FURTHER INFORMATION CONTACT: For technical questions related to the application or requests for an ACE Portal Account, including ACE Protest Filer Accounts, contact the ACE Account Service Desk by calling 1-866-530-4172, selecting option 1, then option 2, or by emailing ACE.Support@cbp.dhs.gov for assistance.

SUPPLEMENTARY INFORMATION:

I. Automated Commercial Environment (ACE)

A. The National Customs Automation Program

The National Customs Automation Program (NCAP) was established by Subtitle B of Title VI—Customs Modernization in the North American Free Trade Agreement (NAFTA) Implementation Act (Customs Modernization Act) (Pub. L. 103-182, 107 Stat. 2057, 2170, December 8, 1993) (19 U.S.C. 1411). Through NCAP, the thrust of customs modernization was on trade compliance and the development of ACE, the planned successor to the Automated Commercial System (ACS). ACE is an automated and electronic system for commercial trade processing which is intended to streamline business processes, facilitate growth in trade, ensure cargo security, and foster participation in global commerce, while ensuring compliance with U.S. laws and regulations and reducing costs for CBP and all of its communities of interest. The ability to meet these objectives depends on successfully modernizing