additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: August 11, 2016.

Michelle Trout,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–19545 Filed 8–16–16; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

National Cancer Institute; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the National Cancer Advisory Board, September 07, 2016, 09:00 a.m. to September 07, 2016, 5:00 p.m., National Institutes of Health, Building 31, 31 Center Drive, Conference Room 10, Bethesda, MD 20892 which was published in the Federal Register on August 08, 2016, 81 FR 52452.

This meeting notice has been amended to change the end time of the open session to 2:45 p.m. The closed session has also been amended to begin at 3:00 p.m. and end at 4:15 p.m. The meeting is partially closed to the public.

Dated: August 11, 2016.

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

[FR Doc. 2016–19544 Filed 8–16–16; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

Prospective Grant of Exclusive Patent License: Development of T Cell Receptors (TCRs) Targeting the KRAS G12D Mutation for the Treatment of Cancer

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: This is notice, in accordance with 35 U.S.C. 209 and 37 CFR part 404, that the National Cancer Institute, National Institutes of Health, Department of Health and Human Services, is contemplating the grant of an exclusive patent license to Kite Pharma, Inc. ("Kite") located in Santa Monica, CA to practice the inventions embodied in the following patent applications:

Intellectual Property


The patent rights in these inventions have been assigned to the government of the United States of America. The prospective exclusive license territory may be worldwide and the field of use may be limited to the development, manufacture and commercialization of retrovirally-engineered mutated KRAS TCR-based autologous peripheral blood T cell therapy products as set forth in the Licensed Patent Rights for the treatment of human cancers.

DATES: Only written comments and/or applications for a license which are received by the National Cancer Institute on or before September 1, 2016 will be considered.

ADDRESSES: Requests for copies of the patent application, inquiries, and comments relating to the contemplated exclusive license should be directed to: Andrew Burke, Ph.D., Licensing and Patenting Manager, Technology Transfer Center, National Cancer Institute, 9609 Medical Center Drive, MSC 9702, Rockville, MD 20852; Telephone: (240) 276–5484; Email andy.burke@nih.gov.

SUPPLEMENTARY INFORMATION: The present invention describes an isolated T cell receptor (TCR) which recognizes the G12D mutation of the Kristen rat sarcoma viral oncogene homolog (KRAS) protein within the context of major histocompatibility complex HLA–A11 presentation.

KRAS is an oncogene with a well-characterized role in the formation of several human cancers, including: Pancreatic, colorectal and lung. Certain mutations, such as the substitution of aspartic acid or valine for glycine at codon 12 (termed G12D and G12V, respectively), occur at relatively high frequency and may represent amenable targets for immunotherapies. Due to the restricted expression of KRAS G12D in pre-cancerous and malignant cells, engineered T cell therapies based on the present invention may be useful for the treatment of select cancers.

The prospective exclusive license will be royalty bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR part 404. The prospective exclusive license may be granted unless within fifteen (15) days from the date of this published notice, the NCI receives written evidence and argument that establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR part 404.

Complete applications for a license in an appropriate field of use that are timely filed in response to this notice will be treated as objections to the grant of the contemplated exclusive license. Comments and objections submitted to this notice will not be made available for public inspection and, to the extent permitted by law, will not be released under the Freedom of Information Act, 5 U.S.C. 552.

Dated: August 8, 2016.

Richard U. Rodriguez,
Associate Director, Technology Transfer Center, National Cancer Institute.

[FR Doc. 2016–19549 Filed 8–16–16; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

National Institute on Aging; 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.


Date: September 29–30, 2016.

Time: 4:00 p.m. to 12:00 p.m.

Agenda: To review and evaluate grant applications.

Place: DoubleTree by Hilton Bethesda, 8120 Wisconsin Ave., Bethesda, MD 20814.

Contact Person: Jeanette L. Johnson, Ph.D., Deputy Review Branch Chief, National Institutes of Health, National Institute on Aging, Gateway Building, Bethesda, MD
DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

30-Day Notice of Proposed Information Collection: Public Housing Agency (PHA) Lease and Grievance Requirements

AGENCY: Office of the Chief Information Officer, HUD.

ACTION: Notice.

SUMMARY: HUD has submitted the proposed information collection requirement described below to the Office of Management and Budget (OMB) for review, in accordance with the Paperwork Reduction Act. The purpose of this notice is to allow for an additional 30 days of public comment.

DATES: Comments Due Date: September 16, 2016.

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB Control Number and should be sent to: HUD Desk Officer, Office of Management and Budget, New Executive Office Building, Washington, DC 20503; fax: 202–395–5806. Email: OIRA_Submission@omb.eop.gov.

FOR FURTHER INFORMATION CONTACT: Colette Pollard, Reports Management Officer, QMAC, Department of Housing and Urban Development, 451 7th Street SW., Washington, DC 20410; email Colette Pollard at Colette.pollard@hud.gov or telephone 202–402–3400. This is not a toll-free number. Persons with hearing or speech impairments may access this number through TTY by calling the toll-free Federal Relay Service at (800) 877–8339.

Copies of available documents submitted to OMB may be obtained from Ms. Pollard.

SUPPLEMENTARY INFORMATION: This notice informs the public that HUD is seeking approval from OMB for the information collection described in Section A.

The Federal Register notice that solicited public comment on the information collection for a period of 60 days was published on March 4, 2016 at 81 FR 11584.

A. Overview of Information Collection

Title of Information Collection: Public Housing Agency (PHA) Lease and Grievance Requirements.

OMB Approval Number: 2577–0006.

Type of Request: Reinstatement of currently approved collection.

Form Number: None.

Description of the need for the information and proposed use: The Public Housing lease and grievance procedures are a recordkeeping requirement on the part of Public Housing agencies (PHAs) as they are required to enter into and maintain lease agreements for each individual or family that occupies a Public Housing unit. Also, both PHAs and tenants are required to follow the protocols set forth in the grievance procedures for both an informal and formal grievance hearing. This information collection is a revision of the previous submission. The reduction in burden hours is attributable to a fewer number of tenants in public housing covered by these lease and grievance procedures.

Respondents (i.e., affected public): Public Housing Authorities (PHAs).

Estimated Number of Respondents: 945,539.

Estimated Number of Responses: 1,359,284.

Frequency of Response: 1.

Average Hours per Response: .25.

Total Estimated Burden: 330,939 hours.

B. Solicitation of Public Comment

This notice is soliciting comments from members of the public and affected parties concerning the collection of information described in Section A on the following:

(1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) The accuracy of the agency’s estimate of the burden of the proposed collection of information;

(3) Ways to enhance the quality, utility, and clarity of the information to be collected; and

(4) Ways to minimize the burden of the collection of information on those who are to respond; including through the use of appropriate automated collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

HUD encourages interested parties to submit comment in response to these questions.


Dated: August 9, 2016.

Colette Pollard,
Department Reports Management Officer, Office of the Chief Information Officer.

[FR Doc. 2016–19546 Filed 8–16–16; 8:45 am]