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 Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, Biomarkers of Autoimmunity in Type 1 Diabetes.

Date: June 14, 2006.

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

Agenda: To review and evaluate grant applications.

Place: Double Tree Rockville, 1750 Rockville Pike, Rockville, MD 20852.

Contact Person: Xiaodu Guo, MD, Ph.D., Scientific Review Administrator, Review Branch, DEA, NIDDK, National Institutes of Health, Room 927, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–4719, *guox@extra.niddk.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.947, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Disease, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: April 30, 2006.

Anna Snouffer,

Acting Director, Office of Federal Advisory Committee Policy.

[FR Doc. 06–4298 Filed 5–8–06; 8:45am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License: The Development of C-6 and C-8 Modified cAMP-Derivatives for the Treatment of Cancer

AGENCY: National Institutes of Health, Public Health Service, HHS.

ACTION: Notice.

SUMMARY: This notice, in accordance with 35 U.S.C. 209(c)(1) and 37 CFR Part 404.7(a)(1)(i), that the National Institutes of Health (NIH), Department of Health and Human Services (HHS), is contemplating the grant of an exclusive patent license to practice the inventions embodied in U.S. Patent Application No. 07/198,489 filed May 23, 1988, entitled ''Use of 8-Cl-cAMP as Anticancer Drug'' [HHS Reference No. E–132–1988/0–US–01], PCT Application filed May 19, 1989 [HHS Reference No. E–132–1988/0–PCT–02], U.S. Patent Application No. 07/896,452 filed June 4, 1992, entitled ''Use of 8-Cl-cAMP as Anticancer Drug'' [HHS Reference No. E–132–1988/0–US–04], U.S. Patent 5,792,752 filed October 27, 1994 and issued August 11, 1998, entitled ''Use of 8-Cl-cAMP as Anticancer Drug'' [HHS Reference No. E–132–1988/0–US–05], U.S. Patent 5,902,794 filed September 22, 1997 and issued May 11, 1999, entitled ''Use of 8-Cl-cAMP as Anticancer Drug'' [HHS Reference No. E–132–1988/0–US–06] and Canadian Patent Application No. 133572 filed May 19, 1989, entitled ''Use of 8-Cl-cAMP as Anticancer Drug'' [HHS Reference No. E–132–1988/0–CA–03], to Kuhnil Pharm. Co. Ltd., which has offices in Seoul, Republic of Korea. The patent rights in these inventions have been assigned and/or exclusively licensed 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 treatment of cancer with 8-Cl-cAMP.

DATES: Only written comments and/or applications for a license which are received by the NIH Office of Technology Transfer on or before June 10, 2006 will be considered.

ADDRESSES: Requests for copies of the patent application, inquiries, comments, and other materials relating to the contemplated exclusive license should be directed to: David A. Lambertson, Ph.D., Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852–3804; Telephone: (301) 435–4632; Facsimile: (301) 402–0220; E-mail: *lambertsond@od.nih.gov.*

SUPPLEMENTARY INFORMATION: Cyclic AMP (cAMP) is a natural biological product with a number of regulatory functions at physiological levels. At higher than physiological concentrations, cAMP has the ability to inhibit the aberrant growth of malignant cells. Because cAMP is a natural product involved in normal biological function, this inhibition occurs without causing significant toxicity. However, this is not a feasible method for treating cancer *in vivo* because of potential interference with the physiological role of cAMP.

C-6 and C-8 modified cAMP derivatives also inhibit the growth of malignant cells. One such derivative, 8-Cl-cAMP, has effectively decreased tumor growth in vitro and in vivo. Specifically, 8-Cl-cAMP showed the ability to decrease tumor growth in leukemia mouse models and xenografts of human tumors. Because of the low toxicity associated with 8-Cl-cAMP, this compound has promise as an anti-cancer agent, particularly with regard to hematological malignancies.

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 404.7. The prospective exclusive license may be granted unless within sixty (60) days from the date of this published notice, the NIH 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 404.7.

Applications for a license in the field of use 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: May 2, 2006.

David R. Sadowski,

Acting Director, Division of Technology Development and Transfer, Office of Technology Transfer, National Institutes of Health.

[FR Doc. E6–6986 Filed 5–8-06; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Citizenship and Immigration Services

Agency Information Collection Activities: Extension of a Currently Approved Information Collection; Comment Request.

ACTION: 30-Day Notice of Information Collection under Review: Application for Certificate of Citizenship, Form N–600. OMB Control No. 1615–0057.

The Department of Homeland Security, U.S. Citizenship and Immigration Services (USCIS) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection was previously published in the **Federal Register** on February 28, 2006, at 71 FR 10048. The notice allowed for a 60-day public comment period. No comments were received on this information collection.

The purpose of this notice is to allow an additional 30 days for public comments. Comments are encouraged and will be accepted until June 8, 2006. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the

estimated public burden and associated response time, should be directed to the Department of Homeland Security (DHS), USCIS, Director, Regulatory Management Division, Clearance Office, 111 Massachusetts Avenue, 3rd floor, Washington, DC 20529. Comments may also be submitted to DHS via facsimile to 202–272–8352 or via e-mail at *rfs.regs@dhs.gov*. When submitting comments by e-mail please make sure to add OMB Control Number 1615–0057 in the subject box. Written comments and suggestions from the public and affected agencies should address one or more of the following four points:

(1) Evaluate 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) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

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

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

Overview of this information collection:

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Application for Certificate of Citizenship.

(3) *Agency form number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection:* Form N–600. U.S. Citizenship and Immigration Services.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract: Primary:* Individuals or households. The information collected on the Form N–600 is necessary for U.S. Citizenship and Immigration Services (USCIS) to make a determination that the citizenship eligibility requirements and conditions are met by the applicant.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 88,500 responses at 1 hour and 35 minutes.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 140,095 annual burden hours.

If you have additional comments, suggestions, or need a copy of the proposed information collection instrument with instructions, or additional information, please visit the USCIS Web site at: *http://uscis.gov/graphics/formsfee/forms/pra/index.htm*.

If additional information is required contact: USCIS, Regulatory Management Division, 111 Massachusetts Avenue, 3rd Floor, Washington, DC 20529 (202) 272–8377.

Dated: May 4, 2006.

Richard A. Sloan,

Director, Regulatory Management Division, U.S. Citizenship and Immigration Services.

[FR Doc. E6–7018 Filed 5–8–06; 8:45 am]

BILLING CODE 4410–10–P

INTER-AMERICAN FOUNDATION

Sunshine Act; Meetings

TIME AND DATE: May 22, 2006, 9 a.m.–12:30 p.m.

PLACE: 901 N. Stuart Street, Tenth Floor, Arlington, Virginia 22203.

STATUS: Open session except for the portion specified as closed session as provided in 22 CFR part 1004.4 (f).

MATTERS TO BE CONSIDERED:

- Personnel Issues.
- Approval of the Minutes of the October 14, 2005, Meeting of the Board of Directors.
- President's Report.
- Fellowship Program.
- RedEAmérica Update.
- Congressional Activities and Strategy.
- Advisory Council.

PORTIONS TO BE OPEN TO THE PUBLIC:

- Approval of the Minutes of the October 14, 2005, Meeting of the Board of Directors.
- President's Report.
- Fellowship Program.
- RedEAmérica Update.
- Congressional Activities and Strategy.
- Advisory Council.

PORTIONS TO BE CLOSED TO THE PUBLIC:

- Personnel issues. Closed session as provided in 22 CFR part 1004.4 (f).

FOR FURTHER INFORMATION CONTACT: Jennifer R. Hodges, General Counsel, (703) 306–4320.

Dated: May 4, 2006.

Jennifer R. Hodges,

General Counsel.

[FR Doc. 06–4342 Filed 5–5–06; 10:30 am]

BILLING CODE 7025–01–P

DEPARTMENT OF THE INTERIOR

Bureau of Reclamation

California Bay-Delta Public Advisory Committee Public Meeting

AGENCY: Bureau of Reclamation, Interior.

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the California Bay-Delta Public Advisory Committee (Committee) will meet on May 25, 2006. The agenda for the Committee meeting will include discussions with State and Federal agency representatives on the status of the revitalization of the CALFED Bay-Delta Program including the 10-Year Action Plan and Program Plans, priority setting, the changing roles of the CALFED agencies, and the restructuring of the Committee Subcommittees.

DATES: The meeting will be held on Thursday, May 25, 2006, from 9 a.m. to 4 p.m. If reasonable accommodation is needed due to a disability, please contact Colleen Kirtlan at (916) 445–5511 or TDD (800) 735–2929 at least 1 week prior to the meeting.

ADDRESSES: These meetings will be held at the John E. Moss Federal Building located at 650 Capitol Mall, 5th Floor, Sacramento, California.

FOR FURTHER INFORMATION CONTACT: Keith Coolidge, California Bay-Delta Authority, at (916) 445–5551, or Diane Buzzard, U.S. Bureau of Reclamation, at (916) 978–5022.

SUPPLEMENTARY INFORMATION: The Committee was established to provide advice and recommendations to the Secretary of the Interior on implementation of the CALFED Bay-Delta Program. The Committee makes recommendations on annual priorities, integration of the eleven Program elements, and overall balancing of the four Program objectives of ecosystem restoration, water quality, levee system integrity, and water supply reliability. The Program is a consortium of State and Federal agencies with the mission to develop and implement a long-term comprehensive plan that will restore ecological health and improve water management for beneficial uses of the San Francisco/Sacramento and San Joaquin Bay Delta.

Committee agendas and meeting materials will be available prior to all meetings on the California Bay-Delta Authority Web site at *http://calwater.ca.gov* and at the meetings. These meetings are open to the public. Oral comments will be accepted from