MSC 7814, Bethesda, MD 20892, (301) 435–1789.

Name of Committee: Center for Scientific Review Special Emphasis Panel.

Date: August 11, 2000. Time: 8:30 a.m. to 12 p.m.

Agenda: To review and evaluate grant applications.

Place: Georgetown Holiday Inn, Mirage 1 Room, 2101 Wisconsin Avenue, NW, Washington, DC 20007.

Contact Person: Prabha L. Ateya, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5152, MSC 7842, Bethesda, MD 20892, (301) 435– 8367.

Name of Committee: Center for Scientific Review Special Emphasis Panel.

Date: August 13–14, 2000. Time: 7:30 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The American Inn, 8130 Wisconsin Ave, Bethesda, MD 20814.

Contact Person: David J. Remondini, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6154, MSC 7890, Bethesda, MD 20892, (301) 435– 1038, remondid@csr.nih.gov

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine, 93.306; 93.333, Clinical Research, 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: July 21, 2000.

### LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 00–19137 Filed 7–27–00; 8:45 am]

BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

Prospective Grant of Exclusive License: "Transcription Factor Decoy and Tumor Growth Inhibitor"

**AGENCY:** National Institutes of Health, Public Health Service, DHHS.

**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, Department of Health and Human Services, is contemplating the grant of an exclusive world-wide license to U.S. Patent Applications 08/977,643, entitled: "Transcription Factor Decoy and Tumor Growth Inhibitor" plus, if available, corresponding foreign patent applications, to Genta Incorporated having a place of business in Lexington, MA. The patent rights in these

inventions have been assigned to the United States of America and the contemplated license may be limited for use in the development and commercialization of diagnostic and therapeutic modalities to treat various diseases and inhibit tumor growth based on the use of cAMP Response Element-palindrome Oligonucleotide as a transcription factor decoy to regulate gene expression (i.e., gene transcription and translation).

**DATES:** Only written comments and/or applications for a license which are received by NIH on or before September 26, 2000 will be considered.

ADDRESSES: Requests for copies of the patent applications, inquiries, comments and other materials relating to the contemplated licenses should be directed to: J. R. Dixon, Ph.D.,
Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville,
Maryland 20852–3804. Telephone: (301) 496–7735 ext. 206; Facsimile: (301) 402–0220. A signed Confidentiality Agreement will be required to receive copies of the patent application.

SUPPLEMENTARY INFORMATION: The technology disclosed in USPA SN: 08/ 977,643 patent application provides compositions and methods for the use of cAMP Response Element-palindrome oligonucleotide as a transcription factor decoy and an inhibitor of tumor growth. Specifically, the 08/977,643 application provides nucleic acid molecules that compete with cAMP Response Elements ("CRE") for binding to transcription factors and a method for regulating gene transcription in target cells comprising: Providing one or more cAMP response element enhancer DNAs and one or more transcription factors that associate with the cAMP response element enhancer DNA; and exposing the target cells to the cAMP response element decoys under condition such that the cAMP response element decoys will compete with the cAMP response element enhancer DNA for binding to the one or more transcription factors.

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

The field of use may be limited to the development and commercialization of

diagnostic and therapeutics modalities to treat various diseases and inhibit tumor growth based on the use of cAMP Response Element-palindrome Oligonucleotide as a transcription factor decoy to regulate gene expression (*i.e.*, gene transcription and translation).

Applications for a license [i.e., completed "Application for License to Public Health Service Inventions"] in the field of use in the development and commercialization of diagnostic and therapeutics modalities to treat various diseases and inhibit tumor growth based on the use of cAMP Response Elementpalindrome Oligonucleotide as a transcription factor decoy to regulate gene expression (i.e., gene transcription and translation) filed in response to this notice will be treated as objections to the grant of the contemplated license. Comments and objections will not be made available for public inspection and, to the extent permitted by law, will not be subject to disclosure under the Freedom of Information Act, 5 U.S.C. 552.

Dated: July 19, 2000.

#### Jack Spiegel,

Director, Division of Technology Development and Transfer, Office of Technology Transfer. [FR Doc. 00–19154 Filed 7–27–00; 8:45 am] BILLING CODE 4140–01–P

# DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-4561-N-47]

Notice of Submission of Proposed Information Collection to OMB; Monthly Report of Excess Income

**AGENCY:** Office of the Chief Information Officer, HUD.

**ACTION:** Notice.

**SUMMARY:** The proposed information collection requirement described below has been submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

**DATES:** Comments Due Date: August 28, 2000.

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB approval number (2502–0086) and should be sent to: Joseph F. Lackey, Jr., OMB Desk Officer, Office of Management and Budget, Room 10235, New Executive Office Building, Washington, DC 20503.