

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 Institute of Child Health and Human Development Special Emphasis Panel Arousal and Attention Regulation in High Risk Children.

*Date:* April 27, 2005.

*Time:* 3:30 p.m. to 6 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, 6100 Executive Boulevard, Room 5B01, Rockville, MD 20852, (telephone conference call).

*Contact Person:* Marita R. Hopmann, PhD, Scientific Review Administrator, Division of Scientific Review, National Institute of Child Health and Human Development, 6100 Building, Room 5B01, Bethesda, MD 20892, (301) 435-6911, [hopmannm@mail.nih.gov](mailto:hopmannm@mail.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. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS).

Dated: April 8, 2005.

**Anna P. Snouffler,**

*Acting Director, Office of Federal Advisory Committee Policy.*

[FR Doc. 05-7547 Filed 4-14-05; 8:45 am]

**BILLING CODE 4140-01-M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Prospective Grant of an Exclusive License: Therapeutics for the Treatment of Retinopathy

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

**ACTION:** Notice.

**SUMMARY:** This notice, in accordance with 35 U.S.C. 209(c)(1) and 37 CFR Part 404.7(a)(1)(i), announces that the National Institutes of Health, Department of Health and Human Services, is contemplating the grant of an exclusive license to practice the inventions embodied in 1. E-223-1992/0-US-01, "SIGNAL TRANSDUCTION INHIBITOR COMPOUNDS", by Elise Kohn, Lance Liotta, Christian Felder, issued Pat No. 5,359,078 (issue date October 25, 1994); 2. E-223-1992/0-US-02, "SIGNAL TRANSDUCTION INHIBITOR TRIAZOLE AND DIAZOLE

COMPOUNDS" by Elise Kohn, Lance Liotta, Christian Felder, issued Pat No. 5,482,954 (issue date January 9, 1996); 3. E-223-1992/0-US-03, "SIGNAL TRANSDUCTION INHIBITOR 1,2,3-TRIAZOLO COMPOUNDS" by Elise Kohn, Lance Liotta, Christian Felder, issued Pat No. 5,498,620, (issued date March 12, 1996); 4. E-223-1992/0-US-04, "SIGNAL TRANSDUCTION INHIBITOR COMPOUNDS" by Elise Kohn, Lance Liotta, Christian Felder, issued Pat No 5,705,514 (issued date January 6, 1998); 5. E-223-1992/0-US-05, "SIGNAL TRANSDUCTION INHIBITOR COMPOUNDS" by Elise Kohn, Lance Liotta, Christian Felder, issued Pat No 5,880,129 (issued date March 9, 1999); 6. E-068-1991/1-US-01, "METHOD FOR INHIBITING METALLOPROTEINASE EXPRESSION" by Elise Kohn, Lance Liotta, issued Pat No. Pat No. 5,602,156 (issued date February 11, 1997); 7. E-220-1993/1-US-01 "METHOD FOR INHIBITING ANGIOGENESIS" by Elise Kohn, Lance Liotta and Riccardo Alessandro issued patent No 5,744,492 (issue date April 28, 1998), to RFE Pharma, having a place of business in Framingham, MA. The patent rights in these inventions have been assigned to the United States of America.

The prospective exclusive license territory may be worldwide, and the field of use may be limited to therapeutics for the treatment of retinopathy.

**DATES:** Only written comments and/or license applications which are received by the National Institutes of Health on or before June 14, 2005, will be considered.

**ADDRESSES:** Requests for copies of the patent and/or patent applications, inquiries, comments and other materials relating to the contemplated exclusive license should be directed to: John Stansberry, Ph.D., Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852-3804; telephone: (301) 435-5236; Facsimile: (301) 402-0220; e-mail: [stansbej@mail.nih.gov](mailto:stansbej@mail.nih.gov).

**SUPPLEMENTARY INFORMATION:** The present invention provides composition of matter claims and methods for inhibiting angiogenesis in a host using carboxyamido-triazole (CAI) and related analogs. The calcium influx inhibitor and matrix metalloproteinase expression inhibitor, CAI, has shown anti-cancer activity due to its ability to influence signal transduction pathways. CAI and CAI analogues inhibit

endothelial cell adhesion and migration in response to basement membrane components and thus block new vessel formation. Pharmaceutical applications directed to inhibiting angiogenesis offer novel approaches to the treatment of cancer, diabetic retinopathy, hemangiomas, vasculidities and other diseases associated with angiogenesis.

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, the NIH receives written evidence and argument that establish 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: April 6, 2005.

**Steven M. Ferguson,**

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

[FR Doc. 05-7542 Filed 4-14-05; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Prospective Grant of Exclusive License: Methods for Treating Inflammatory Bowel Disease Using Cholera Toxin B Subunit

**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 patent license to practice the inventions embodied in U.S. Patent Application 10/129,907, filed May 10, 2002 [DHHS Ref. E-263-1999/0-US-03], entitled "Methods for treating inflammatory bowel disease using cholera toxin B subunit," to SBL Vaccin AB, which is located in Stockholm, Sweden. The patent rights in these inventions have

been assigned to the United States of America.

The prospective exclusive license territory may be worldwide and the field of use may be limited to the use of cholera toxin B as a therapeutic treatment of inflammatory bowel disease, specifically Crohn's disease.

**DATES:** Only written comments and/or applications for a license which are received by the NIH Office of Technology Transfer on or before June 14, 2005, 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: Michelle A. Booden, Ph.D., Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852-3804; telephone: (301) 451-7337; Facsimile: (301) 402-0220; e-mail: [boodenm@mail.nih.gov](mailto:boodenm@mail.nih.gov).

**SUPPLEMENTARY INFORMATION:** The technology describes a method of treating or preventing inflammatory bowel disease by administering cholera toxin B subunit (CT-B). Specifically, the patent application discloses administering CT-B as a method for treating and preventing Crohn's disease (CD) and Ulcerative Colitis (UC) as well as a method for treating and preventing inflammation and/or autoimmune disorders mediated by increased interferon gamma (INF- $\gamma$ ) and or interleukin 12 (IL-12).

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, 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: April 6, 2005.

**Steven M. Ferguson,**

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

[FR Doc. 05-7543 Filed 4-14-05; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-4980-N-15]

### Federal Property Suitable as Facilities To Assist the Homeless

**AGENCY:** Office of the Assistant Secretary for Community Planning and Development, HUD.

**ACTION:** Notice.

**SUMMARY:** This Notice identifies unutilized, underutilized, excess, and surplus Federal property reviewed by HUD for suitability for possible use to assist the homeless.

**FOR FURTHER INFORMATION CONTACT:** Kathy Ezzell, room 7266, Department of Housing and Urban Development, 451 Seventh Street SW., Washington, DC 20410; telephone (202) 708-1234; TTY number for the hearing- and speech-impaired (202) 708-2565 (these telephone numbers are not toll-free), or call the toll-free Title V information line at 1-800-927-7588.

**SUPPLEMENTARY INFORMATION:** In accordance with 24 CFR part 581 and section 501 of the Stewart B. McKinney Homeless Assistance Act (42 U.S.C. 11411), as amended, HUD is publishing this Notice to identify Federal buildings and other real property that HUD has reviewed for suitability for use to assist the homeless. The properties were reviewed using information provided to HUD by Federal landholding agencies regarding unutilized and underutilized buildings and real property controlled by such agencies or by GSA regarding its inventory of excess or surplus Federal property. This Notice is also published in order to comply with the December 12, 1988 Court Order in *National Coalition for the Homeless v. Veterans Administration*, No. 88-2503-OG (D.D.C.).

Properties reviewed are listed in this Notice according to the following categories: Suitable/available, suitable/unavailable, suitable/to be excess, and unsuitable. The properties listed in the three suitable categories have been reviewed by the landholding agencies, and each agency has transmitted to HUD: (1) Its intention to make the property available for use to assist the homeless, (2) its intention to declare the property excess to the agency's needs, or (3) a statement of the reasons that the property cannot be declared excess or made available for use as facilities to assist the homeless.

Properties listed as suitable/available will be available exclusively for homeless use for a period of 60 days from the date of this notice. Where

property is described as for "off-site use only" recipients of the property will be required to relocate the building to their own site at their own expense.

Homeless assistance providers interested in any such property should send a written expression of interest to HHS, addressed to Heather Ranson, Division of Property Management, Program Support Center, HHS, room 5B-17, 5600 Fishers Lane, Rockville, MD 20857; (301) 443-2265. (This is not a toll-free number.) HHS will mail to the interested provider an application packet, which will include instructions for completing the application. In order to maximize the opportunity to utilize a suitable property, providers should submit their written expressions of interest as soon as possible. For complete details concerning the processing of applications, the reader is encouraged to refer to the interim rule governing this program, 24 CFR part 581.

For properties listed as suitable/to be excess, that property may, if subsequently accepted as excess by GSA, be made available for use by the homeless in accordance with applicable law, subject to screening for other Federal use. At the appropriate time, HUD will publish the property in a notice showing it as either suitable/available or suitable/unavailable.

For properties listed as suitable/unavailable, the landholding agency has decided that the property cannot be declared excess or made available for use to assist the homeless, and the property will not be available.

Properties listed as unsuitable will not be made available for any other purpose for 20 days from the date of this notice. Homeless assistance providers interested in a review by HUD of the determination of unsuitability should call the toll free information line at 1-800-927-7588 for detailed instructions or write a letter to Mark Johnston at the address listed at the beginning of this notice. Included in the request for review should be the property address (including zip code), the date of publication in the **Federal Register**, the landholding agency, and the property number.

For more information regarding particular properties identified in this notice (*i.e.*, acreage, floor plan, existing sanitary facilities, exact street address), providers should contact the appropriate landholding agencies at the following addresses: **Energy:** Mr. Andy Duran, Department of Energy, Office of Engineering & Construction Management, ME-90, 1000 Independence Ave, SW., Washington, DC 20585; (202) 586-4548; **GSA:** Mr.