

session will be 8:00 a.m. until 9:30 a.m. The meeting is partially closed to the public.

Dated: August 31, 2001.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 01-22785 Filed 9-10-01; 8:45 am]

BILLING CODE 4140-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), 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 552(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 Institute of Allergy and Infectious Diseases Special Emphasis Panel.

Date: September 21, 2001.

Time: 1:00 pm to 5:00 pm.

Agenda: To review and evaluate grant applications.

Place: 6700B Rockledge Drive, Bethesda, MD 20892-2616, (Telephone Conference Call).

Contact Person: Nasrin Nabavi, PhD, Scientific Review Administrator, Scientific Review Program, Division of Extramural Activities, NIAID, NIH, Room 2156, 6700B Rockledge Drive, MSC 7610, Bethesda, MD 20892-7610, 301 496-2550, nn301@nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: August 31, 2001.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 01-22786 Filed 9-10-01; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), 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: National Institute of Allergy and Infectious Diseases Special Emphasis Panel.

Date: September 21, 2001.

Time: 9:00 am to 11:00 am.

Agenda: To review and evaluate grant applications.

Place: 6700B Rockledge Drive, Bethesda, MD 20892-2616, (Telephone Conference Call).

Contact Person: Nasrin Nabavi, PhD, Scientific Review Administrator, Scientific Review Program, Division of Extramural Activities, NIAID, NIH, Room 2156, 6700B Rockledge Drive, MSC 7610, Bethesda, MD 20892-7610, 301-496-2550, nn301@nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: August 31, 2001.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 01-22787 Filed 9-10-01; 8:45 am]

BILLING CODE 4140-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License: Virus-Like Particles for the Induction of Autoantibodies

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 (NIH), Department of Health and Human Services, is contemplating the grant of an exclusive license to practice the invention embodied in: United States Patent Application 09/835,124 and its foreign equivalents entitled "Virus-Like Particles for the Induction of Autoantibodies" filed on April 13, 2001, with priority back to U.S. S/N 60/105,132, filed October 21, 1998 to Cytos Biotechnology AG, having a place of business in Zurich, Switzerland. The patent rights in this invention have been assigned to the United States of America.

DATES: Only written comments and/or application for a license which are received by the NIH Office of Technology Transfer on or before November 13, 2001 will be considered.

ADDRESSES: Requests for a copy of the patent application, inquiries, comments and other materials relating to the contemplated license should be directed to: Peter Soukas, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852-3804; Email: ps193c@nih.gov; Telephone: (301) 496-7056, ext. 268; Facsimile: (301) 402-0220.

SUPPLEMENTARY INFORMATION: This invention claims compositions and methods for producing antibodies to tolerogens (self-antigens normally exposed to B cells that fail to induce an antibody response.) The compositions of the invention comprise multiple copies of a tolerogen (or at least one B cell epitope of a tolerogen) chimerized to viral virus-like particle in an orderly manner. This invention could potentially replace any treatment utilizing chronic administration of a monoclonal antibody that reacts with a self-antigen.

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 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 alphavirus virus-like particle (VLP), Hepatitis B Virus (HBV)-capsid protein VLP, and bacteriophage Q β VLP, and Ty VLP vaccines.

Properly filed competing applications for a license filed in response to this notice will be treated as objections to the contemplated license. Comments

and objections submitted in response 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: September 4, 2001.

Jack Spiegel,

Director, Division of Technology Development and Transfer, Office of Technology Transfer.

[FR Doc. 01-22790 Filed 9-10-01; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License: Veterinary Vaccine Against Escherichia Coli O157 Infection Composed of Detoxified LPS Conjugated to Proteins

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

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 (NIH), Department of Health and Human Services, is contemplating the grant of an exclusive license to practice the invention embodied in: United States Patent Application 09/744,289 and its foreign equivalents entitled "Vaccine Against Escherichia Coli O157 Infection Composed of Detoxified LPS Conjugated to Proteins" filed January 22, 2001, with priority back to PCT/US98/14976, filed July 20, 1998 to Fort Dodge Animal Health, a Division of American Home Products, having a place of business in Overland Park, Kansas. The patent rights in this invention have been assigned to the United States of America.

DATES: Only written comments and/or application for a license which are received by the NIH Office of Technology Transfer on or before November 13, 2001 will be considered.

ADDRESSES: Requests for a copy of the patent application, inquiries, comments and other materials relating to the contemplated license should be directed to: Peter Soukas, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852-3804; Email: ps193c@nih.gov; Telephone: (301) 496-7056, ext. 268; Facsimile: (301) 402-0220.

SUPPLEMENTARY INFORMATION: This invention comprises the O-specific

polysaccharide of Shiga toxin-producing bacteria, particularly *E. coli* O157, conjugated to a carrier protein such as *Pseudomonas aeruginosa* recombinant exoprotein A or hepatitis B surface or core antigen. This vaccine is suitable for use in humans and animals. Cattle are carriers of *E. coli* O157, and are the primary reservoir of *E. coli* O157 by shedding the bacteria into the environment. Fifty percent (50%) of cattle are estimated to be carriers of *E. coli* O157. Use of this vaccine in cattle could eliminate *E. coli* O157 and prevent contamination of meat in slaughterhouses.

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 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 *E. coli* conjugate vaccines for veterinary use.

Properly filed competing applications for a license filed in response to this notice will be treated as objections to the contemplated license. Comments and objections submitted in response 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: September 4, 2001.

Jack Spiegel, Ph.D.,

Director, Division of Technology Development and Transfer, Office of Technology Transfer.

[FR Doc. 01-22792 Filed 9-10-01; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License: Adeno-Associated Virus with Inverted Terminal Repeat Sequence as Promoter

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

ACTION: Notice.

SUMMARY: This is notice in accordance with 15 U.S.C. 209(c)(1) and 37 CFR 404.7(a)(1)(i) that the National Institutes of Health (NIH), Department of Health and Human Services, is contemplating the grant of an exclusive world-wide license to practice the inventions

embodied in any U.S. patents 5,587,308 (12/24/1996); 5,989,540 (11/23/1999); 5,866,696 (02/02/1999), and 6,165,781 (12/26/2000) or foreign applications corresponding to PCT Patent Application PCT/US93/05310, entitled "Modified Adeno-Associated Virus Vector Capable of Expression from a Novel Promoter" published as WO 93/24641 (12/09/1993) to Targeted Genetics Corporation of Seattle, Washington. The prospective exclusive license may be limited to the development of compositions and methods utilizing Adeno-Associated Viral Vectors which are useful in the treatment and prophylaxis of human and animal diseases.

DATES: Only written comments and/or applications for a license which are received by NIH on or before November 13, 2001, will be considered.

ADDRESSES: Requests for a copy of these patent applications, inquiries, comment and other materials relating to the contemplated license should be directed to Susan S. Rucker, J.D., Patent and Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852-3804; telephone: 301/496-7056 ext 245; fax: 301/402-0220. A signed Confidentiality Agreement will be required to receive copies of the patent applications.

SUPPLEMENTARY INFORMATION: The patents describe and claim compositions and methods utilizing adeno-associated viral (AAV) vectors. In particular, these vectors utilize the AAV Inverted Terminal Repeat (ITR) as the promoter element to control expression of the nucleic acid encoding the heterologous protein to be delivered to the patient. The ability of these vectors to utilize the AAV ITR as the promoter increases the capacity of the AAV vector with respect to the size of the heterologous protein which can be encoded and delivered via the vector. The methods of the patent can be used to deliver and produce therapeutic or prophylactic products which are particularly useful in the field of gene therapy.

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. This 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.