

Control and Prevention (CDC), Technology Transfer Office, Department of Health and Human Services (DHHS), is contemplating the grant of a limited field of use, exclusive license in India to practice the inventions embodied in the patent referred to below to Molecular Diagnostic Laboratory, having a place of business in Lucknow, India. The patent rights in these inventions have been assigned to the government of the United States of America. The patent(s) to be licensed are:

US 6,896,892 B2 entitled "Insecticide-Impregnated Fabric and Method of Production," issue date 05.24.2005. CDC Technology ID No. I-008-99.

*Status:* Issued.

*Issue Date:* 05.24.2005

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.

*Technology:* This technology provides a new insecticide-impregnated fabric and method of production for bednets.

**SUPPLEMENTARY INFORMATION:** In accordance with 35 U.S.C. 209(e) and 37 CFR 404.7(a)(1)(i), CDC is providing public notice of its intention to grant an exclusive license. CDC will accept written comments concerning this notice for 30 days. Applications for a license filed in response to this notice will be treated as objections to the grant of the contemplated license. Only written comments and/or applications for a license which are received by CDC within thirty days of this notice will be considered. 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.

**ADDRESSES:** Requests for a copy of this patent, inquiries, comments, and other materials relating to the contemplated license should be directed to Suzanne Seavello Shope, J.D., Technology Licensing and Marketing Specialist, Technology Transfer Office, Centers for Disease Control and Prevention (CDC), 4770 Buford Highway, Mailstop K-79, Atlanta, GA 30341, telephone: (770) 488-8613; facsimile: (770) 488-8615.

Dated: August 31, 2006.

**James D. Seligman,**  
Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E6-14871 Filed 9-7-06; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Prospective Grant of Exclusive License: Diagnostics of Fungal Infections

**AGENCY:** Technology Transfer Office, Centers for Disease Control and Prevention (CDC), Department of Health and Human Services.

**ACTION:** Notice.

**SUMMARY:** This is a notice in accordance with 35 U.S.C. 209(e) and 37 CFR 404.7(a)(1)(i) that the Centers for Disease Control and Prevention (CDC), Technology Transfer Office, Department of Health and Human Services (DHHS), is contemplating the grant of a worldwide, limited field of use, co-exclusive license to practice the inventions embodied in the patent and patent applications referred to below to Myconostica, Inc. (Myconostica) having a place of business in Manchester, United Kingdom. CDC intends to grant no more than three licenses to these inventions. The patent rights in these inventions have been assigned to the government of the United States of America. The patent and patent applications to be licensed are:

*Title:* Nucleic Acids for Detecting *Aspergillus* Species and Other Filamentous Fungi.

*U.S. Patent Application Serial No.:* 09/423,233.

*Filing Date:* 6/27/2000.

*Domestic Status:* 6,372,430.

*Issue Date:* 4/16/2002.

*Title:* Molecular Identification of *Aspergillus* Species.

*U.S. Patent Application Serial No.:* 60/381,463.

*Filing Date:* 5/17/2002.

*Domestic Status:* Pending.

*Issue Date:* N/A.

*Title:* Nucleic Acids for the Identification of Fungi and Methods for Using the Same.

*U.S. Patent Application Serial No.:* 60/325,241.

*Filing Date:* 9/26/2001.

*Domestic Status:* Pending.

*Issue Date:* N/A.

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.

Specific DNA (oligonucleotide) probes have been developed for a wide variety of systemic disease causing fungi, including *Aspergillus* species and others. A probe has been developed for identification of all dimorphic fungi.

These probes can be used for the rapid identification of fungal pathogens and for the diagnosis of mycotic diseases.

**ADDRESSES:** Requests for a copy of these patent applications, inquiries, comments, and other materials relating to the contemplated license should be directed to Andrew Watkins, Director, Technology Transfer Office, Centers for Disease Control and Prevention (CDC), 4770 Buford Highway, Mailstop K-79, Atlanta, GA 30341, telephone: (770) 488-8610; facsimile: (770) 488-8615. Applications for a license filed in response to this notice will be treated as objections to the grant of the contemplated license. Only written comments and/or applications for a license which are received by CDC within sixty days of this notice will be considered. 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. A signed Confidential Disclosure Agreement will be required to receive a copy of any pending patent application.

Dated: August 31, 2006.

**James D. Seligman,**  
Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E6-14872 Filed 9-7-06; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. 2006N-0349]

#### Risk Communication on Medical Devices: Sharing Perspectives

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of public meeting.

**SUMMARY:** The Food and Drug Administration (FDA), in cooperation with the Advanced Medical Technology Association (AdvaMed), is announcing a public meeting entitled "Risk Communication on Medical Devices: Sharing Perspectives." This 1-day workshop is intended to bring together various creators and recipients of medical device risk/benefit information to discuss how this information is developed, disseminated, and perceived; and to explore ways in which the process might be improved.

**DATES AND TIMES:** The public meeting will be held on September 26, 2006, from 7:30 a.m. to 5 p.m. Online registration is available until 5 p.m. on