

Institutes of Health, Department of Health and Human Services, is contemplating the grant of an exclusive license to practice the inventions embodied in U.S. Patent Application S/N 08/533,895, filed on September 26, 1995, entitled "MHC Class II Restricted Melanoma Antigens and Their Use in Therapeutic Methods", to Therion Biologics Corporation of Cambridge, Massachusetts. The patent rights in these inventions have been assigned to the United States of America.

The prospective exclusive license territory will be worldwide and the field of use may be limited to recombinant poxvirus-based vaccines for human cancer immunotherapy, said poxviruses encoding Class II-restricted melanoma antigens, or modifications, derivatives, or immunogenic peptides thereof, and vaccination protocols comprising the administration of one or more Class II-restricted melanoma peptides in addition to a recombinant poxvirus-based vaccine (for example, in a prime and boost protocol), but specifically excluding the use of these peptides in any context other than a recombinant poxvirus-based vaccination protocol.

**DATES:** Only written comments and/or license applications which are received by the National Institutes of Health on or before June 29, 2001 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: Elaine White, M.B.A., Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD. 20852-3804. Telephone: (301) 496-7056, X282; Facsimile (301) 402-0220; E-mail eg46t@nih.gov.

**SUPPLEMENTARY INFORMATION:** 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 23, 2001.

**Jack Spiegel,**

*Director, Division of Technology Development and Transfer, Office of Technology Transfer.*  
[FR Doc. 01-10578 Filed 4-27-01; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Prospective Grant of Exclusive License: Human Papilloma Inhibition by Antisense Oligonucleotides

**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: Korean Patent Application 10-2000-7002392 entitled "Human Papilloma Inhibition by Antisense Oligonucleotides" filed on June 30, 2000, to Gyn-Gen Bio, Inc., having a place of business in Seoul, Korea. 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 June 29, 2001 will be considered.

**ADDRESSES:** Requests for a copy of the patent applications, 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:** The present invention relates to the use of antisense oligonucleotides to inhibit Human Papilloma Virus (HPV). The antisense oligonucleotides have a phosphorothioate backbone structure and sequences complementary to portions of the human papilloma virus 16 E6 gene. See the equivalent United States patent number 6,084,090 and Alvarez-Salas et al., "Growth inhibition of cervical tumor cells by antisense oligodeoxynucleotides directed to the

human papillomavirus type 16 E6 gene," *Antisense Nucleic Acid Drug Dev* 1999 Oct;9(5):441-50 for further details.

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 treatment and prevention of Human Papilloma Virus infection with antisense oligonucleotides. The licensed territory is expected to be limited to Korea, China, Malaysia and Thailand.

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: April 20, 2001.

**Jack Spiegel,**

*Director, Division of Technology Development and Transfer, Office of Technology Transfer*  
[FR Doc. 01-10577 Filed 4-27-01; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Prospective Grant of Exclusive License: Identification of TRP-2 as a New Human Tumor Antigen Recognized by Cytotoxic T Lymphocytes

**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 license to practice the inventions embodied in U.S. Patent Applications S/N 08/725,736, filed on October 4, 1996, and now U.S. Patent 5,831,016 which issued on November 3, 1998; S/N 09/161,877 (DIV of 08/725,736), filed on September 28, 1998, and now U.S. Patent 6,132,980 which issued on October 17, 2000; S/N 09/162,368 (DIV of 08/725,736), filed on September 28,