

DEPARTMENT OF HEALTH AND HUMAN SERVICES**National Institutes of Health****Government-Owned Inventions; Availability for Licensing**

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

ACTION: Notice.

SUMMARY: The inventions listed below are owned by agencies of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 207 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

ADDRESSES: Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852-3804; telephone: 301/496-7057; fax: 301/402-0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

Vibrio cholerae O139 Conjugate Vaccines

Shousun Szu, Zuzana Kossaczka, John Robbins (NICHD)

DHHS Reference No. E-274-00/1; PCT/US00/24119 filed 01 Sep 2000

Licensing Contact: Peter Soukas; 301/496-7056 ext. 268; e-mail: soukasp@od.nih.gov

Cholera remains an important public health problem. Epidemic cholera is caused by two *Vibrio cholerae* serotypes O1 and O139. The disease is spread through contaminated water. According to information reported to the World Health Organization in 1999, nearly 8,500 people died and another 223,000 were sickened with cholera worldwide. This invention is a polysaccharide-protein conjugate vaccine to prevent and treat infection by *Vibrio cholerae* O139 comprising the capsular polysaccharide (CPS) of *V. cholerae* O139 conjugated through a dicarboxylic acid dihydrazide linker to a mutant diphtheria toxin carrier. In addition to the conjugation methods, also claimed in the invention are methods of immunization against *V. cholerae* O139 using the conjugates of the invention. The inventors have shown that the conjugates of the invention elicited in

mice high levels of serum antibodies to CPS, a surface antigen of *Vibrio cholerae* O139, that have vibriocidal activity. Clinical trials of the two most immunogenic conjugates have been planned by the inventors. This invention is further described in *Infection and Immunity* 68(9), 5037-5043, Sept. 2000.

Inhibition of MXR Transport by Acridine Derivatives

Susan Bates, Robert Robey (NCI)
DHHS Reference No. E-258-99/0 filed 20 Jan 2000

Licensing Contact: Vasant Gandhi; 301/496-7056 ext. 224; e-mail: gandhiv@od.nih.gov

The invention relates to a new use for a compound, an acridine derivative, as an inhibitor of multidrug resistance in cancer cells. Specifically, the inventors have shown that the compound modulates the transport of compounds from mitoxantrone-resistant (MXR) cells wherein the cells overexpress an MXR gene. The MXR gene is also known by the following designations: BCRP, ABCP, and ABCG2.

Jack Spiegel,

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

[FR Doc. 00-32815 Filed 12-22-00; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES**National Institutes of Health****Prospective Grant of Exclusive License: Hydroxylamine Compositions for the Prevention or Retardation of Cataracts**

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 404.7(a)(1)(i), that the National Institutes of Health (NIH), Department of Health and Human Services, in contemplating the grant of an exclusive license worldwide to practice the invention embodied in: U.S. Patent Number 6,001,853 issued December 14, 1999 entitled, "Hydroxylamine Compositions for the Prevention or Retardation of Cataracts", to SL Pharmaceuticals, having a place of business in Kennett Square, PA 19348. The contemplated exclusive license may be limited to use for human therapeutics and diagnostics. The United States of

America is the assignee of the patent rights in this invention.

DATES: Only written comments and/or application for a license which is received by the NIH Office of Technology Transfer on or before February 26, 2001, will be considered.

ADDRESSES: Request for a copy of the patent, inquiries, comments and other materials relating to the contemplated license should be directed to: Marlene Shinn, Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852-3804; Telephone: 301-496-7056 ext. 285; Facsimile: 301-402-0220; E-mail: ms482m@nih.gov.

SUPPLEMENTARY INFORMATION: Cataracts are believed to be a disease of multifactorial origin involving many of the same processes that characterize the process of aging in other issues. This invention relates generally to a pharmaceutical composition and treatment to inhibit the development of cataracts in the crystalline lens of the eye by administering a hydroxylamine to a patient at risk of developing a cataract. This technology is an improvement over what is presently known, in that it allows for a clinically useful non-surgical treatment that retards the development of age-related cataracts.

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.

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: December 15, 2000.

Jack Spiegel,

Director, Division of Technology Development and Transfer, Office of Technology Transfer.
[FR Doc. 00-32816 Filed 12-22-00; 8:45 am]

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