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#### **Imaging With Positron-Emitting Taxanes as a Guide to Antitumor Therapy**

Jerry M. Collins, Raymond W. Klecker, Lawrence Anderson (FDA)  
Serial No. 60/155,061 filed 21 Sep 1999

The present application discloses the use of positron-emitting compounds to label taxane type drugs. This invention also describes methods of synthesizing these taxane type compounds. Further, methods to guide treatment of solid tumors, with labeled taxanes, are also disclosed in the present application. Advantages of using this technology include: (1) Avoidance of exposing patients to toxic drugs that have no potential for benefit; (2) ability to rapidly determine whether a given tumor will be likely to respond to a particular drug; and (3) the ability to monitor the impact of various dosages, schedules, and modulators for delivery, *in situ*, at the actual tumor under treatment conditions.

#### **Conjugate Vaccine for *Neisseria Meningitidis***

Xin-Xing Gu (NIDCD) and Chao-Ming Tsai (FDA)  
Serial No. 60/148,021 filed 10 Aug 1999

The invention discloses a vaccine which comprises lipooligosaccharide (LOS) isolated from *N. meningitidis* and conjugated to a carrier protein. The invention also discloses a method of making the acellular vaccine. The method consists of two main steps. In the first step the lipooligosaccharide (LOS), chosen so it does not contain the lacto-N-neotetraose human antigen (LNnT), is detoxified by a novel procedure which uses hydrazine to remove the O-linked fatty acids. In the second step, the detoxified LOS (dLOS) is covalently conjugated to a carrier protein such as Tetanus Toxoid (TT). The dLOS produced in step 1 is 10,000 fold less toxic than the parent LOS. The conjugate vaccine exhibited a high level of immunogenicity as evidenced by the high titer of IgG antibody to native LOS, obtained in mice and rabbits. The rabbit antisera produced by the conjugate vaccine of one *N. meningitidis* strain (strain 7880, A,L10) exhibited bactericidal activity and cross reactivity with heterologous *N. meningitidis* strains. A conjugate vaccine made in this method may be multivalent, composed of dLOSs from different strains and/or immunotypes of *N. meningitidis* and will thus protect against all types of *N. meningitidis*, including type B.

A portion of this invention was disclosed in a poster by Tsai, Gu and Quakyi at the Fifth Conference of the International Endotoxin Society held in Santa Fe, New Mexico in September 12–15, 1998.

Dated: April 7, 2000.

#### **Jack Spiegel,**

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

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### **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 contacting John Peter Kim, at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852–3804; telephone: 301/496–7056 ext. 264; fax: 301/402–0220; e-mail: jk141n@nih.gov. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

#### **High Speed Parallel Nucleic Acid Sequencing**

Thomas D. Schneider, Denise Rubens (NCI)  
Serial No. 60/151,580 filed 30 Aug 1999

The present application describes a new method and apparatus for DNA sequencing called Two Dye Sequencing (TDS). This method employs engineered DNA polymerases which are labeled with a fluorophore such as Green Fluorescent Protein (GFP) and are combined with an annealed oligonucleotide primer in a chamber of a microscope field of view capable of detecting individual molecules. Four

nucleotide triphosphates, each labeled on the base with a different fluorescent dye are introduced to the reaction. Light of a specific wavelength is used to excite the fluorophore on the polymerase, which in turn excites the neighboring fluorophore on the nucleotide by Fluorescence Resonance Energy Transfer (FRET). As nucleotides are added to the primer, their spectral emissions provide sequence information of the DNA molecule.

#### **Hydrazide Inhibitors of HIV–1 Integrase**

Yves Pommier, Nouri Neamati, Zhaiwai Lin, Terrence R. Burke, Jr. (NCI)  
DHHS Reference Nos. E–037–99/0 filed 12 Mar 1999 and E–037–99/1 filed 10 Mar 2000

The human immunodeficiency virus (HIV) is the causative agent of acquired immunodeficiency syndrome (AIDS). Drug-resistance is a critical factor contributing to the gradual loss of clinical benefit to treatments for HIV infection. Accordingly, combination therapies have further evolved to address the mutating resistance of HIV. However, there has been great concern regarding the apparent growing resistance of HIV strains to current therapies.

It has been found that a certain class of compounds including salicylhydrazides and analogs and derivatives thereof are effective and selective anti-integrase inhibitors which are active in the presence of both Mn(+2) and Mg(+2) and which may be used in the treatment or prevention of infection by HIV and AIDS. The subject invention provides for such compounds and for methods of inhibiting HIV integrase.

Dated: April 5, 2000.

#### **Jack Spiegel,**

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

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### **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

#### **National Institutes of Health**

#### **“Conference on Challenges in Health Disparity in the New Millennium: A Call to Action”**

Notice is hereby given of the NIH Office of Research on Minority Health (ORMH) Conference on Challenges in Health Disparity in the New Millennium: A Call to Action, which will be held April 16–19, 2000, at the