

and Use of Information Collection: A case-control study will examine the relationship between exposure to benzene and the risk of lymphohematopoietic malignancies and related disorders and lung cancer in Chinese workers. Cases and controls will be selected from participants in a recent cohort study of benzene-exposed workers in China. The data will be used by the NCI to examine risk among workers exposed to low levels of benzene, and to characterize the dose and time-specific relationship between benzene exposure and disease risk. Frequency of Response: One-time study. Affected Public: Individuals or households. Type of Respondents: Workers. The annual reporting burden is as follows: Estimated Number of Respondents: 1,545; Estimated Number of Responses per Respondent: One; Average Burden Hours per Response: 0.75; and Estimated Total Annual Burden Hours Requested: 386.

There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

*Request for Comments:* Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection or information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

**FOR FURTHER INFORMATION:** To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Dr. Richard Hayes, Project Officer, National Cancer Institute, Executive Plaza South, Room 8114, Rockville, Maryland 20892-7364, or call non-toll-free number (301) 496-9093, or FAX your request to (301) 402-1819, or E-mail your request, including your address, to [HayesR@exchange.nih.gov](mailto:HayesR@exchange.nih.gov).

**COMMENTS DUE DATE:** Comments regarding this information collection are best assured of having their full effect if received on or before February 26, 2001.

Dated: December 18, 2000.

**Reesa L. Nichols,**

*NCI Project Clearance Liaison.*

[FR Doc. 00-33085 Filed 12-27-00; 8:45 am]

**BILLING CODE 4140-01-M**

## 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.

#### Transgenic Zebrafish with Vascular Specific Expression of Exogenous Genes Driven by the Zebrafish Fli-1 Promoter

Brant M. Weinstein, Nathan N. Lawson (NICHD)

DHHS Reference No. E-003-01/0

*Licensing Contact:* Marlene Shinn; 301/496-7056 ext. 285; email: [shinnm@od.nih.gov](mailto:shinnm@od.nih.gov)

The technology portrayed in this invention is available through a Biological Materials License for research tools and diagnostic tests. Zebrafish are an important and valuable model system for high-throughput mutational or pharmacological screens for genes or molecules with important roles in blood vessel growth or differentiation. This invention consists of germline transgenic zebrafish lines in which the expression of green fluorescent protein (EGFP) is driven by zebrafish Fli-1 promoter sequences. These transgenic lines display bright, uniform, and persistent expression of EGFP protein

throughout the vascular system. The Fli promoter also drives transient EGFP expression in cranial neural crest and its derivatives. The transgenics allow straightforward, noninvasive fluorescent visualization of virtually all blood vessels in the animal throughout embryonic and early larval development.

These Fli-EGFP transgenics have a number of potential applications. They can be used to help identify endogenous genes important for blood vessel formation, either by screening mutagenized transgenic embryos for vascular specific mutants or by preparing vascular specific cDNA libraries for use in novel gene discovery. They also provide an efficient method for performing high-throughput in vivo screening for antiangiogenic or proangiogenic drugs and other compounds. Using transgenic zebrafish for these screens has the added benefit of simultaneously revealing toxic and teratogenic effects of the tested agents on a whole, developing organism.

#### Transcranial Magnetic Stimulation Coil for Specific Non-Invasive Deep Brain Stimulation

Abraham Zangen (NIDA), Roy Wise (NIDA), Mark Hallett (NINDS), Yiftach Roth (EM), Pedro Miranda (NINDS)

DHHS Reference No. E-223-00/0 filed 20 Oct 2000

*Licensing Contact:* Dale Berkley; 301/496-7735 ext. 223; e-mail: [berkleyd@od.nih.gov](mailto:berkleyd@od.nih.gov)

The invention is a magnetic stimulator that is placed in contact with the head of a subject to magnetically stimulate the brain. The invention has applications in the treatment of neurophysiological or cardiovascular conditions, and may be of particular utility in the treatment of disorders associated with deep regions of the brain, such as drug addiction and depression. The unique coil shape of the stimulator is designed to target deep brain regions like the nucleus accumbens, which are associated with the biological mechanism underlying drug abuse. Deep regions of the brain are also implicated in depressive disorders, and this coil is likely to offer an improvement in the transcranial magnetic stimulation therapy currently being tested for treatment of depression.

#### Peroxynitrite Generators, Compositions Comprising Same, and Methods for Treating Biological Disorders Using Same

Challice L. Bonifant, Joseph E. Saavedra and Larry K. Keefer (NCI)

DHHS Reference No. E-175-00/0 filed 02 June 2000

*Licensing Contact:* Norbert Pontzer; 301/496-7735, ext. 284; e-mail: pontzern@od.nih.gov

Diazoniumdiolates are a class of compounds which release nitric oxide (NO) under physiological conditions. Nitric oxide performs a number of regulatory functions in vivo such as controlling vascular tone and platelet function, but it can also combine with superoxide ion to produce peroxynitrite ion, as especially reactive species. Peroxynitrite-mediated cellular toxicity may have several therapeutic applications. Because of the relatively low amounts of superoxide ion present in some cells, the peroxynitrite mechanism of diazoniumdiolate toxicity is not uniformly available. In order to generate peroxynitrite ions in tissues or other media lacking adequate levels of superoxide ion, this invention provides a new class of compounds which release NO and superoxide ion simultaneously to generate peroxynitrite ions.

Molecules of this invention can be designed to generate peroxynitrite ion at specific biochemical targets. For one type of targeting, the release of NO is designed to be triggered by nucleophilic attack on the diazoniumdiolate drug while superoxide generation is simultaneously occurring at a quinone moiety elsewhere in the molecule. If the required nucleophilic attack is designed to be specifically catalyzed in the active site of glutathione S-transferase-pi, a cytoprotective enzyme overexpressed by certain tumors to render them drug-resistant, compounds of this invention could restore the susceptibility of tumor cells to chemotherapy by knocking out the excess enzyme, thereby preventing the tumor cells from inactivating the chemotherapeutic agents. Attachment of the compounds to polymeric compositions would physically localize the peroxynitrite activity. Physical localization in vivo may have utility against the recently recognized chronic infections caused by biofilms, and generation of peroxynitrite ions in vitro may have utility against infectious bilfilms on medical devices.

Dated: December 20, 2000.

**Jack Spiegel,**

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

[FR Doc. 00-33086 Filed 12-27-00; 8:45 am]

**BILLING CODE 4140-01-M**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Heart, Lung, and Blood Institute; 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 Heart, Lung, and Blood Institute Special Emphasis Panel Program Project Reviews.

*Date:* January 12, 2001.

*Time:* 8 AM to 2 PM.

*Agenda:* To review and evaluate grant applications.

*Place:* Sheraton Columbia Hotel, 10207 Wincopin Circle, Columbia, MD 21044.

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*Contact Person:* Jeffrey H. Hurst, PhD, Health Scientist Administrator, 6701 Rockledge Drive, Room 7208, Bethesda, MD 20892, 301-435-0303.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: December 20, 2000.

**LaVerne Y. Stringfield,**

*Director, Office of Federal Advisory Committee Policy.*

[FR Doc. 00-33084 Filed 12-27-00; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Institute of Neurological Disorders and Stroke; Notice of 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 a meeting of the Board of Scientific Counselors, National Institute of Neurological Disorders and Stroke.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the National Institute of Neurological Disorders and Stroke, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Board of Scientific Counselors, National Institute of Neurological Disorders and Stroke

*Date:* February 4-6, 2001.

*Closed:* February 4, 2001, 7:00 PM to 10:00 PM.

*Agenda:* To review and evaluate personal qualifications and performance, and competence of individual investigators.

*Place:* Bethesda Marriott Hotel, 5151 Pooks Hill Road, Bethesda, MD 20814.

*Open:* February 5, 2001, 8:15 AM to 11:10 AM.

*Agenda:* To discuss program planning and project accomplishments.

*Place:* National Institutes of Health, Natcher Building, Conference Room F-1/2, Bethesda, MD 20892.

*Closed:* February 5, 2001, 11:10 AM to 1:15 PM.

*Agenda:* To review and evaluate personal qualifications and performance, and competence of individual investigators.

*Place:* National Institutes of Health, Natcher Building, Conference Room F-1/2, Bethesda, MD 20892.

*Open:* February 5, 2001, 1:15 PM to 4:15 PM.

*Agenda:* To discuss program planning and program accomplishments.

*Place:* National Institutes of Health, Natcher Building, Conference Room F-1/2, Bethesda, MD 20892.

*Closed:* February 5, 2001, 4:15 PM to 5:15 PM.

*Agenda:* To review and evaluate personal qualifications and performance, and competence of individual investigators.

*Place:* National Institutes of Health, Natcher Building, Conference Room F-1/2, Bethesda, MD 20892.

*Closed:* February 5, 2001, 6:00 PM to 10:00 PM.