This invention is a vaccine for the prevention of disease caused by nontypeable *H. influenzae*, which causes 25-40% of otitis media cases (middle ear infections) in children and other respiratory tract diseases in humans. The emergence of antibioticresistant bacteria has caused concern that treatment of otitis media will become more problematic. This invention offers a new approach to managing otitis media. The vaccine is composed of lipooligosaccharide, isolated from the surface of strains of nontypeable H. influenzae and treated with hydrazine to remove esterified fatty acids, covalently conjugated to an immunogenci carrier, such as tetanus toxoid. The conjugates have been shown to be nontoxic by the limulus amebocyte assay, rabbit pyrogen test, and in a mouse lethal toxicity test. Antisera raised in rabbits immunized with the conjugate is bactericidal. (portfolio: Infectious Diseases—Vaccines, bacterial)

Materials And Methods for Detection and Treatment of Insulin Dependent Diabetes

NK Maclaren, AL Notkins, Q Li, MS Lan (NIDR)

Serial No. 08/514,213 filed 11 Aug 95 and

Serial No. 08/548,159 filed 25 Oct 95 Licensing Contact: J. Peter Kim, 301/ 496–7056 ext. 264

Insulin-dependent diabetes mellitus (IDDM) affects close to one million people in the United States. It is an autoimmune disease in which the immune system produces antibodies that attack the body's own insulinmanufacturing cells in the pancreas. Patients require daily injections of insulin to regulate blood sugar levels. The invention identified two proteins, named IA-2 and IA-2β, that are important markers for type I (juvenile, insulin-dependent) diabetes. IA-2/IA-2β, when used in diagnostic tests, recognized autoantibodies in 70 percent of IDDM patients. Combining IA-2/IA-2β with other known markers increased the level of identification to 90 percent of individuals with IDDM. Moreover, the presence of autoantibodies to IA-2/ IA-2β in otherwise normal individuals was highly predictive in identifying those at risk of ultimately developing clinical disease. It is now possible to develop a rapid and effective test that can screen large populations for IDDM. In addition, IĂ–2/IÅ–2β are candidates for immune tolerance and prevention of disease development.

Compositions Comprising Vitamin F C Weinberger, S Kitareewan (NIEHS) Serial No. 60/003,443 filed 08 Sep 95; PCT/US96/15205 filed 06 Sep 96 Licensing Contact: Carol Lavrich, 301/ 496–7056 ext. 287

This invention relates to a collection of potential fat-soluble vitamins that may coordinate animal metabolism and development. RXR is a nuclear receptor that plays a central role in cell signaling by heterodimerizing with receptors binding thyroid hormones, retinoids and vitamin D. The invention and others of its compositions can be characterized as likely physiological effectors that may represent essential components for human nutrition and cell growth. Thus, the invention suggests that it may coordinate cell physiology through RXR-dependent hormone signaling pathways.

Macrocyclic Chelates, And Methods of Use Thereof

OA Gansow, K Kumar (NCI) Serial No. 08/140,714 filed 22 Oct 93 U.S. Patent 5,428,154 issued 27 Jul 95 Licensing Contact: Raphe Kantor, 301/ 496–7735 ext. 247

Substituted 1,4,7,10-tetraaza cyclododecane-N,N', N", N"'-tetraacetic acid (DOTA) has numerous desirable chelating qualities that make it useful for treating a number of cdellular disorders. Presently available chelating agents lack specificity for their intended targets or do not adequately bind the chelated metal ion. These substituted DOTAs have a strong affinity for a number of metal ions. They can also be linked to biomolecules to form systems for delivering the chelated metal ion, which can be radiolabeled, to specific sites within a cell or organelle. (portfolio: Cancer—Therapeutics, immunoconjugates, conjugate chemistry)

The Cloning of Perilipin Proteins

C Londos, AS Greenberg, AR Kimmel, JJ Egan (NIDDK) Serial No. 08/132,649 filed 04 Oct 93 U.S. Patent 5,585,462 to issue 17 Dec 96 Licensing Contact: Ken Hemby, 301/ 496/7735 ext. 265

Perilipins are found at the surface of lipid storage droplets of adipocytes. Little is known about the molecules on the surface of lipid droplets that may be involved in lipid metabolism and trafficking. The present invention provides isolated nucleic acid sequences which encode a family of perilipin proteins as well as isolated, purified perilipin proteins. These are useful as markers for differentiation of true adipocyte cells from non-adipocyte cells which, as a result of pathophysiological conditions, assume

adipocyte characteristics. (portfolio: Cancer—Research Materials)

Dated: December 6, 1996. Barbara M. McGarey, Deputy Director, Office of Technology Transfer.

[FR Doc. 96–31883 Filed 12–16–96; 8:45 am] BILLING CODE 4140–01–M

## National Cancer Institute, Notice of Meeting

Notice is hereby given of the meeting of the National Cancer Institute Board of Scientific Advisors Prevention Working Group, January 30–31, 1997 at the Crystal Gateway Marriott, 1700 Jefferson Davis Highway, Arlington, Virginia.

This meeting will be closed to the public on January 30-31, 1997 from 8:30 a.m. to approximately 10 p.m. each day for the discussion of confidential issues relating to the review, discussion and evaluation of individual programs and projects conducted by the NCI Prevention Program. These discussions will reveal confidential trade secrets or commercial property such as patentable material, and personal information including consideration of personnel qualifications and performance, the competence of individual investigators and similar matters, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Information pertaining to the meeting may be obtained from Dr. Jack Gruber, Executive Secretary, National Cancer Institute Prevention Working Group, National Cancer Institute, 6130 Executive Blvd., EPN, Rm. 540, Bethesda, MD 20892 (301–496–9740).

Dated: December 10, 1996.

Paula N. Hayes,

Acting Committee Management Officer, NIH. [FR Doc. 96–31879 Filed 12–16–96; 8:45 am] BILLING CODE 4140–01–M

# National Cancer Institutes; 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 of the National Cancer Institute Special Emphasis Panel (SEP):

Name of SEP: Cooperative Family Registry for Epidemiologic Studies of Colon Cancer Date: January 9–10, 1997

Time: 9:00 am

Place: Executive Plaza North, Room G 6130 Executive Boulevard Bethesda, MD 20852 Contact Person: Lalita D. Palekar, Ph.D. Scientific Review Administrator National Cancer Institute, NIH Executive Plaza North, Room 643 6130 Executive Boulevard, MSC 7405 Bethesda, MD 20892–7405 Telephone: 301/496–7575

Purpose/Agenda: To evaluate and review responses to RFA CA96–011, entitled Cooperative Family Registry for Epidemiologic Studies of Colon Cancer.

The meeting will be closed in accordance with the provisions set forth in secs. 552b(c)(4) and 552b(c)(6), Title 5, U.S.C. Applications and the discussions could reveal confidential trade secrets or commercial property such as patentable material and personal information concerning individuals associated with the applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

(Catalog of Federal Domestic Assistance Program Numbers: 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control)

Dated: December 10, 1996.

Paula N. Hayes,

Acting Committee Management Officer, NIH. [FR Doc. 96–31880 Filed 12–16–96; 8:45 am] BILLING CODE 4140–01–M

# National Cancer Institutes; 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 of the National Cancer Institute Special Emphasis Panel (SEP):

Name of SEP: Neuropsychological Testing for Children and Adults with Chronic Medical Illness (Cancer and HIV–1 infection). Date: January 8, 1997.

Time: 9:00 am.

Place: Executive Plaza North, Room H, 6130 Executive Boulevard, Bethesda, MD 20852.

Contact Person: Lalita D. Palekar, Ph.D., Scientific Review Administrator, National Cancer Institute, NIH Executive Plaza North, Room 643, 6130 Executive Boulevard, MSC 7405, Bethesda, MD 20892–7405, Telephone: 301/496–7575.

*Purpose/Agenda:* To evaluate and review responses to NO1 SC 71004–09.

The meeting will be closed in accordance with the provisions set forth in secs. 552(c)(4) and 552(c)(6), Title 5, U.S.C. Applications and the discussions could reveal confidential trade secrets or commercial property such as patentable material and personal information concerning individuals associated with the applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

(Catalog of Federal Domestic Assistance Program Numbers: 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control)

Dated: December 10, 1996.

Paula N. Hayes,

Acting Committee Management Officer, NIH. [FR Doc. 96–31881 Filed 12–16–96; 8:45 am] BILLING CODE 4140–01–M

#### National Institute of Diabetes and Digestive and Kidney Diseases; 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:

Committee Name: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel (Telephone Conference Call).

Date: December 27, 1996.

Time: 1:00 p.m.

*Place*: Natcher Building, Room 6AS–25F, National Institutes of Health, 45 Center Drive, Bethesda, Maryland 20892–6600.

Contact Person: Lakshmanan Sankaran, Ph.D., Scientific Review Administrator, Natcher Building, Room 6AS–25F, National Institutes of Health, 45 Center Drive, Bethesda, Maryland 20892–6600, Phone: 301–594–7799.

*Agenda/Purpose:* To review and evaluate a research grant application.

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

The meeting will be closed in accordance with the provisions set forth in secs. 552b(c)(4) and 552b(c)(6), Title 5, U.S.C. Applications and/or proposals and the discussions could reveal confidential trade secrets or commercial property such as patentable material and personal information concerning individuals associated with the applications and/or proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

(Catalog of Federal Domestic Assistance Program No. 93.847–849, Diabetes, Endocrine and Metabolic Diseases; Digestive Diseases and Nutrition; and Kidney Diseases, Urology and Hematology Research, National Institutes of Health)

Dated: December 11, 1996.

Paula N. Hayes,

Acting Committee Management Officer, NIH. [FR Doc. 96–31976 Filed 12–16–96; 8:45 am] BILLING CODE 4140–01–M

## DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-3967-N-03]

### Notice of Proposed Information Collection for Public Comment

**AGENCY:** Office of the Assistant Secretary for Public and Indian Housing, HUD.

**ACTION:** Notice.

SUMMARY: The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

**DATES:** Comments due: February 18, 1997

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB Control Number and should be sent to: Mildred M. Hamman, Reports Liaison Officer, Public and Indian Housing, Department of Housing and Urban Development, 451—7th Street, SW, Room 4240, Washington, D.C. 20410–5000.

FOR FURTHER INFORMATION CONTACT: Mildred M. Hamman, (202) 708–0846, (This is not a toll-free number.) for copies of the proposed forms and other available documents.

SUPPLEMENTARY INFORMATION: The Department will submit the proposed information collection to OMB for review, as required by the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35, as amended).

This Notice is soliciting comments from members of the public and affecting agencies concerning the proposed collection of information to: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond; including through the use of appropriate automated collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

This Notice also lists the following information: