

desiring to make formal oral presentations should notify the contact person before February 1, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: January 10, 2002.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 02-1409 Filed 1-18-02; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed collection; Comment Request; Ethical Issues Associated With Nurse Practitioner and Physician Assistant Practice: A Comparative Analysis

SUMMARY: In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the Department of Clinical Bioethics, the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

PROPOSED COLLECTION: *Title:* Ethical Issues Associated with Nurse Practitioner and Physician Assistant Practice: A Comparative Analysis. *Type of Information Collection Request:* NEW. *Need and Use of Information Collection:* The purposes of the study are (1) to examine whether the current practice environment has created ethical concerns/conflict for Nurse Practitioners and Physician Assistants in the provision of patient care; (2) to explore relationships between selected individual, organizational, and state regulatory factors and ethical conflict in practice and the perceived delivery of quality care; and (3) to examine the perceived level of ethics preparedness and confidence in ethics decision-making. The findings will provide valuable information concerning: (1) The importance of ethics and ethical factors from the perspective of different professional groups; and (2) ethics educational needs of Nurse Practitioners and Physician Assistants. *Frequency of Response:* Once. *Affected Public:* Individuals; Academic Institutions;

Business or other for-profit; Not-for-profit organizations. *Type of Respondents:* Nurse Practitioners and Physician Assistants. The annual report burden is as follows: *Estimated Number of Respondents:* 1,400; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours Per Response:* .33; and *Estimated Total Annual Burden Hours Requested:* 462. The annualized cost to respondents is estimated at \$33,600. There are no Capital Costs to report. There are no Operating 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 is 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) minimize the burden 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 CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Connie Ulrich, RN, PhD, Principal Investigator, Department of Clinical Bioethics, Warren G. Magnuson Clinical Center, Building 10, Room 1C118, Bethesda, MD 20892, or call non-toll-free number (301) 451-8338 or E-mail your request, including your address to: culrich@cc.nih.gov.

COMMENTS DUE DATE: Comments regarding this information collection are best assured of having their full effect if received within 60-days of the date of this publication.

Dated: January 14, 2002.

David K. Henderson,

Deputy Director, Warren G. Magnuson Clinical Center, National Institutes of Health.

Ezekiel Emanuel,

Director, Department of Clinical Bioethics, Warren G. Magnuson Clinical Center, National Institutes of Health.

[FR Doc. 02-1435 Filed 1-18-02; 8:45 am]

BILLING CODE 4140-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; California Health Interview Survey—Complementary and Alternative Medicine

SUMMARY: In compliance with the requirement of section 2506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

PROPOSED COLLECTION: *Title:* California Health Interview Survey—Complementary and Alternative Medicine (CHIS-CAM). *Type of Information Collection Request:* New. *Need and Use of Information Collection.* The NCI has sponsored a Cancer Control Topical Module (CCTM) to the California Health Interview Survey (CHIS), administered in 2001. The CHIS is a telephone survey designed to provide population-based, standardized health-related data. Initiated by the UCLA Center for Health Policy Research, California Department of Health Services, and the Public Health Institute, the survey was unfunded by a number of public and private sources.

The 2001 CHIS CCTM was similar in content to the 2000 National Health Interview Survey (NHIS) CCTM and was administered to one sample adult in more than 54,000 households. NCI anticipates comparing the CHIS and NHIS data in order to conduct comparative and pooled analyses that will enable better estimates of health-related behaviors and cancer risk factor for smaller racial/ethnic minority populations.

The CHIS-CAM is a cross-sectional telephone survey nested in the CHIS study population of all adult respondents who agreed to be re-contacted. Complementary and Alternative Medicine (CAM) is a rapidly growing component of prevention and treatment of chronic illness in the United States. Yet the study of cancer has been largely excluded from the existing population-based surveys on CAM due to sample size restrictions, and little reliable information exists on how CAM utilization varies among different ethnic groups and among those with chronic illnesses.

The CHIS-CAM survey will be administered to approximately 2,000

cancer survivors and 6,000 non-cancer adults. It will enable NCI to collect extensive information on CAM, cancer and other chronic illnesses, and link it with the breadth of basic data already collected from the large, racially and ethnically diverse sample of CHIS respondents.

Comprehensive and detailed collection of information on CAM will enable NCI to increase its understanding of how, why, and to what effect CAM is used. The CHIS-CAM survey data

will allow NCI to compare individuals who report various types of cancer and other chronic conditions and to determine: (1) The major categories of CAM procedures being used, as well as the specific therapies targeted toward cancer prevention and treatment, (2) how various subgroups in the population (defined by race/ethnicity, gender, age, health status, etc.) compare with regards to CAM procedures being used; (3) to what extent persons with cancer used specific types of CAM

before or after diagnoses with cancer, and whether cancer patients used CAM in place of, or in addition to, conventional medical care; (4) whether systematic CAM treatments for cancer might lead to harm or interact with conventional treatments for cancer; and (5) what expenditures people are paying out-of-pocket for CAM procedures. Frequency of Response: One-time. Affected public: Individuals. Types of respondents: U.S. adults. The annual reporting burden is as follows:

TABLE A—ANNUALIZED BURDEN ESTIMATES FOR CHIS-CAM DATA COLLECTION

Type of respondents	Estimated number of respondents	Estimated no. of responses per respondent	Average burden hours per response	Estimated total annual burden hour requested
U.S. Adults	8,000	1	.35	2,800

There is no annualized cost to respondents. There are no Capital Costs to report. There are no Operating 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 of information is necessary for the proper performance of the functions 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 CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Anita Ambs, Project Coordinator, National Cancer Institute, EPN 4106, 6130 Executive Boulevard, Bethesda Maryland 20892-7344, or call non-toll free number (301) 451-8500 or email your request, including your address to ambsa@mail.nih.gov.

COMMENTS DUE DATE: Comments regarding this information collection are best assured of having their full effect if received within 60 days from the date of this publication.

Dated: January 10, 2002.

Reesa L. Nichols,

NCI Project Clearance Liaison.

[FR Doc. 02-1436 Filed 1-18-02; 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 contacting Marlene Shinn, J.D., at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852-3804; telephone: 301/496-7056 ext. 285; fax: 301/402-0220; e-mail: shinnm@od.nih.gov. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

Thermostable DNA Polymerases that Bypass Lesions in DNA

Dr. Roger Woodgate (NICHD) and Dr. Francois Boudsocq (NICHD)
DHHS Reference No. E-232-01/0—
Research tool

Lesions in DNA often block DNA polymerases, especially in those polymerases used in the Polymerase Chain Reaction (PCR). Old DNA, such as that from forensic samples, is often damaged and cannot be used for PCR analysis.

The NIH announces the identification of two novel Y-family DNA polymerases—called Dbh and Dpo4—from the archaea *Sulfolobus solfataricus* P1 and *Sulfolobus solfataricus* P2, respectively. The Y family of polymerases are characterized by their ability to replicate through DNA lesions that may block the activity of other, more conventional, polymerases such as the thermostable enzymes used in PCR. Both Dbh and Dpo4 enzymes have been shown to be as thermostable as the Taq polymerase (Dpo4, in particular) and can copy stretches of DNA up to 1300 bp in length. Because these polymerases are in general more efficient at coping with DNA lesions, they may be useful in the amplification of damaged DNA and could be useful in forensic PCR applications.

A Novel Human DNA Polymerase, POL IOTA, Involved in DNA Repair and Mutagenesis

Drs. Roger Woodgate and John McDonald (NICHD)
DHHS Reference No. E-229-01/0—
Research tool

The NIH announces the identification of a novel DNA polymerase called POL IOTA, that is highly error prone and