

Announcement and presentation of winners will be at GSA's National Travel Forum 2006 (June 26–29, 2006 in Los Angeles, CA).

Dated: January 6, 2006.

Patrick F. McConnell,

Acting Director, Travel Management Policy.

[FR Doc. E6–168 Filed 1–11–06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Meeting of the President's Council on Bioethics

AGENCY: The President's Council on Bioethics, HHS.

ACTION: Notice.

SUMMARY: The President's Council on Bioethics (Edmund D. Pellegrino, MD, Chairman) will hold its twenty-third meeting, at which, among other things, it will continue the discussion on ethical issues relating to children. Subjects discussed at past Council meetings (though not on the agenda for the present one) include: Cloning, assisted reproduction, reproductive genetics, IVF, ICSI, PGD, sex selection, inheritable genetic modification, patentability of human organisms, neuroscience, aging retardation, lifespan-extension, and organ procurement for transplantation. Publications issued by the Council to date include: Human Cloning and Human Dignity: An Ethical Inquiry (July 2002); Beyond Therapy: Biotechnology and the Pursuit of Happiness (October 2003); Being Human: Readings from the President's Council on Bioethics (December 2003); Monitoring Stem Cell Research (January 2004), Reproduction and Responsibility: The Regulation of New Biotechnologies (March 2004), Alternative Sources of Human Pluripotent Stem Cells: A White Paper (May 2005), and Taking Care: Ethical Caregiving in Our Aging Society (September 2005).

DATES: The meeting will take place Thursday, February 2, 2006, from 9 a.m. to 5:15 p.m. e.t. and Friday, February 3, 2006, from 8:30 a.m. to 12:30 p.m. e.t.

ADDRESSES: The Madison, 15th and M Streets, NW., Washington, DC 20005. Phone 202–862–1600.

Agenda: The meeting agenda will be posted at <http://www.bioethics.gov>.

Public Comments: The Council encourages public input, either in person or in writing. At this meeting, interested members of the public may address the Council, beginning at 11:30 a.m., on Friday, February 3. Comments

are limited to no more than five minutes per speaker or organization. As a courtesy, please inform Ms. Diane Gianelli, Director of Communications, in advance of your intention to make a public statement, and give your name and affiliation. To submit a written statement, mail or e-mail it to Ms. Gianelli at one of the addresses given below.

FOR FURTHER INFORMATION CONTACT: Ms. Diane Gianelli, Director of Communications, The President's Council on Bioethics, Suite 700, 1801 Pennsylvania Avenue, Washington, DC 20006. Telephone: 202/296–4669. E-mail: info@bioethics.gov. Web site: <http://www.bioethics.gov>.

Dated: January 6, 2006.

F. Daniel Davis,

Executive Director, The President's Council on Bioethics.

[FR Doc. 06–276 Filed 1–11–06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–06–0278]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–4766 and send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques

or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

National Hospital Ambulatory Medical Care Survey (NHAMCS) 2007–2008 [OMB No. 0920–0278]—Extension—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The National Hospital Ambulatory Medical Care Survey (NHAMCS) has been conducted annually since 1992. The purpose of NHAMCS is to meet the needs and demands for statistical information about the provision of ambulatory medical care services in the United States. Ambulatory services are rendered in a wide variety of settings, including physicians' offices and hospital outpatient and emergency departments. The target universe of the NHAMCS is in-person visits made to outpatient departments (OPDs) and emergency departments (EDs) of non-Federal, short-stay hospitals (hospitals with an average length of stay of less than 30 days) or those whose specialty is general (medical or surgical) or children's general.

NHAMCS was initiated to complement the National Ambulatory Medical Care Survey (NAMCS, OMB No. 0920–0234) which provides similar data concerning patient visits to physicians' offices. NAMCS and NHAMCS are the principal sources of data on approximately 90 percent of ambulatory care provided in the United States.

NHAMCS provides a range of baseline data on the characteristics of the users and providers of ambulatory medical care. Data collected include patients' demographic characteristics, reason(s) for visit, physicians' diagnosis(es), diagnostic services, medications, and disposition. These data, together with trend data, may be used to monitor the effects of change in the health care system, for the planning of health services, improving medical education, determining health care work force needs, and assessing the health status of the population. In addition, a Cervical Cancer Screening Supplement (CCSS) will be added to collect information on cervical cancer screening practices from hospital OPD clinics. It will allow the CDC/National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP) to evaluate cervical cancer screening methods and the use of HPV tests.