

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.

Users of NHAMCS data include, but are not limited to, congressional offices, Federal agencies, state and local governments, schools of public health, colleges and universities, private

industry, nonprofit foundations, professional associations, clinicians, researchers, administrators, and health planners. NCHS is seeking OMB approval to extend this survey for an

additional three years. There are no costs to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN TABLE

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hours)	Total burden (in hours)
Hospital induction	490	1	55/60	449
ED induction	400	1	1	400
OPD induction	250	4	1	1,000
ED Patient record form	400	100	5/60	3,333
OPD Patient record form	250	200	5/60	4,167
CCSS	250	1	15/60	63
Total				9,412

Dated: January 5, 2006.

Joan F. Karr,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E6-210 Filed 1-11-06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-06-0234]

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 e-mail 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 Ambulatory Medical Care Survey (NAMCS) 2007-2008 (OMB No. 0920-0234)—Extension—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

The NAMCS was conducted annually from 1973 to 1981, again in 1985, and resumed as an annual survey in 1989. The purpose of NAMCS 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 NAMCS target population consists of all office visits made by ambulatory patients to non-Federal office-based physicians (excluding those in the specialties of anesthesiology, radiology, and pathology) who are engaged in direct patient care. For the first time in 2006, physicians and mid-level providers (*i.e.*, nurse practitioners, physician assistants, and nurse midwives) practicing in

community health centers (CHCs) were added to the NAMCS sample, and these data will continue to be collected in 2007-2008. To complement NAMCS data, NCHS initiated the National Hospital Ambulatory Medical Care Survey (NHAMCS, OMB No. 0920-0278) to provide data concerning patient visits to hospital outpatient and emergency departments.

The NAMCS provides a range of baseline data on the characteristics of the users and providers of ambulatory medical care. Data collected include the patients' demographic characteristics, reason(s) for visit, physicians' diagnosis(es), diagnostic services, medications, and visit disposition. In addition, a Cervical Cancer Screening Supplement (CCSS) will continue to be a key focus in 2007-2008. The CCSS collects information on cervical cancer screening practices performed by selected physician specialties. It will allow the CDC/National Center for Chronic Disease Prevention and Health Promotion to evaluate cervical cancer screening methods and the use of human papillomavirus tests.

Users of NAMCS data include, but are not limited to, congressional offices, Federal agencies, state and local governments, schools of public health, colleges and universities, private industry, nonprofit foundations, professional associations, clinicians, researchers, administrators, and health planners. NCHS is seeking OMB approval to extend this survey for an additional three years. There are no costs to the respondents other than their time.