obtained from Marjorie S. Greenberg, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 2402, Hyattsville, Maryland 20782, telephone (301) 458–4245. Information also is available on the NCVHS home page of the HHS Web site: http://www.ncvhs.hhs.gov/, where further information including an agenda will be posted when available. Should you require reasonable accommodation, please contact the CDC Office of Equal Employment Opportunity on (301) 458–4EE0 (4336) as soon as possible.


James Scanlon,
Deputy Assistant Secretary for Planning and Evaluation, Office of the Assistant Secretary for Planning and Evaluation.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics: Meeting

Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) announces the following advisory committee meeting.

Name: National Committee on Vital and Health Statistics (NCVHS), Full Committee Meeting.

Time and Date: March 1, 2012; 9:00 a.m.–3:45 p.m. EST, March 2, 2012; 10:00 a.m.–4:00 p.m. EST.

Place: Double Tree Hilton Hotel Silver Spring, 8727 Colesville Road, Silver Spring, Maryland 20910, Tel: 1–301–589–5200.

Status: Open.

Purpose: At this meeting the Committee will hear presentations and hold discussions on several health data policy topics. On the morning of the first day the Committee will hear updates from the Department (HHS), the Centers for Medicare and Medicaid Services (CMS), and the Office of the National Coordinator (ONC). There will also be discussion on items for approval: (1) Recommendation letter on standards for Claims Attachments; (2) recommendation letter on ACA requirements to seek input on improving standardization and uniformity in new financial and administrative activities beyond those already being addressed under HIPAA; and (3) a recommendation letter on the Standards/Operating Rule Maintenance Process. After lunch, there will be a review of the final action items discussed on the first day. Additionally, the Committee will hear subcommittee reports, strategic plans and discuss next steps. After lunch, there will be a briefing on the new CMS Line of Service for Information Resources Initiative. The public will be invited to comment on the information presented. Further information will be provided on the NCVHS Web site at http://www.ncvhs.hhs.gov/.

The times shown above are for the full Committee meeting. Subcommittee breakout sessions are scheduled for late in the afternoon on the first day and in the morning prior to the full Committee meeting on the second day. Agendas for these breakout sessions will be posted on the NCVHS Web site (URL below) when available. Contact Person for More Information: Substantive program information as well as summaries of meetings and a roster of committee members may be obtained from Marjorie S. Greenberg, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 2402, Hyattsville, Maryland 20782, telephone (301) 458–4245. Information also is available on the NCVHS home page of the HHS Web site: http://www.ncvhs.hhs.gov/, where further information including an agenda will be posted when available. Should you require reasonable accommodation, please contact the CDC Office of Equal Employment Opportunity on (301) 458–4EE0 (4336) as soon as possible.


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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention
[60 Day—12–0814]

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 404–639–7570 or send comments to Kimberly Lane, CDC, Reports Clearance Officer, 1600 Clifton Road, MS D–74, Atlanta, GA 30333 or send an email to obm@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

CDC Cervical Cancer Study (CX3)(OMB No. 0920–0814, exp. 6/30/2012)—Revision—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC). Background and Brief Description

The National Breast and Cervical Cancer Early Detection Program (NBCCEDP) is the only organized national screening program in the United States. The program offers breast and cervical cancer screening to underserved women. Given resource limitations, the screening standards for cervical cancer in the program include an annual Pap test until a woman has had three consecutive normal Pap tests, at which time the Pap test frequency is reduced to every three years. HPV DNA testing has been approved in the U.S. as a secondary screening tool for ASCUS (Atypical Squamous Cells of Undetermined Significance), and as a primary screening tool for women 30 years of age and older, but it is not currently a reimbursable expense under program guidelines. Adopting HPV testing along with Pap testing in women over 30 could help the program better utilize resources by extending the screening interval of women who are cytology negative and HPV test negative, which is estimated to be 80–90% of women. In 2005, the NBCCEDP convened an expert panel to determine policies on reimbursement of the HPV DNA test with the Pap test (co-test) for primary screening. The panel recommended that the program not reimburse for the HPV DNA test but instead requested that pilot studies be performed to measure the feasibility, acceptability and barriers to use of the test.

A pilot study, the CDC Cervical Cancer Study (CX3), is currently being conducted in 15 clinics in the state of Illinois. A total of 2,246 women between the ages of 30 and 60 who visited one