ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090. Please bring photo ID for entry to a Federal building.

FOR FURTHER INFORMATION CONTACT: http://www.hhs.gov/healthit/ahic/ healthrecords/.

SUPPLEMENTARY INFORMATION: The Workgroup will continue its discussion on ways to achieve widespread adoption of certified EHRs, minimizing gaps in adoption among providers.

The meeting will be available via Web cast. For additional information, go to: http://www.hhs.gov/healthit/ahic/ healthrecords/ehr_instruct.html.

Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. 07–6135 Filed 12–21–07; 8:45 am] BILLING CODE 4150–45–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; American Health Information Community Electronic Health Records Workgroup Meeting

ACTION: Announcement of meeting.

SUMMARY: This notice announces the 21st meeting of the American Health Information Community Chronic Care Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. 92–463, 5 U.S.C., App.)

DATES: January 17, 2008, from 1 p.m. to 4 p.m., Eastern Time.

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090. Please bring photo ID for entry to a Federal building.

FOR FURTHER INFORMATION CONTACT:

http://www.hhs.gov/healthit/ahic/ chroniccare/

SUPPLEMENTARY INFORMATION: The Workgroup will hear testimony on ways to use information technology to better coordinate care for patients with chronic conditions and will discuss this information in light of opportunities to better facilitate patient care coordination.

The meeting will be available via Web cast. For additional information, go to:

http://www.hhs.gov/healthit/ahic/ chrnoccare/cc_instruct.html.

Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. 07–6136 Filed 12–21–07; 8:45 am]

BILLING CODE 4150-45-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; American Health Information Community Population Health and Clinical Care Connections Workgroup Meeting

ACTION: Announcement of meeting.

SUMMARY: This notice announces the 22nd meeting of the American Health Information Community Population Health and Clinical Care Connections Workgroup in accordance with the Federal Advisory Committee act (Pub. L. 92–463, 5 U.S.C., App.)

DATES: January 3, 2008, from 1 p.m. to 4 p.m. [Eastern Time].

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090 (please bring photo ID for entry to a Federal building).

FOR FURTHER INFORMATION CONTACT: *http://www.hhs.gov/healthit/ahic/ population/.*

SUPPLEMENTARY INFORMATION: The Workgroup will continue its discussion on how to facilitate the flow of reliable health information among population health and clinical care systems necessary to protect and improve the public's health.

The meeting will be available via Web cast. For additional information, go to: http://www.hhs.gov/healthit/ahic/ population/pop_instruct.html.

Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. 07–6157 Filed 12–21–07; 8:45 am] BILLING CODE 4150–45–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day-08-07AW]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an email to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project

Experimental and Theoretical Study of Early Detection and Isolation of Influenza—New—The National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Federal Occupational Safety and Health Act of 1970, section 501, enables NIOSH to carry out research relevant to the health and safety of workers. Some diseases like influenza and Severe Acute Respiratory Syndrome (SARS) can be spread when people produce clouds of droplets (called aerosols) by coughing or sneezing. Aerosol transmission of infectious diseases is of particular interest today because of increased concern over a possible global influenza pandemic. The possible airborne spread of influenza is especially important to health-care workers and emergency responders, who face a much greater risk of exposure than does the general public. However, substantial gaps exist in our understanding of the generation and spread of infectious aerosols containing influenza. This lack of information hampers the ability of health scientists to model and predict the transmission of influenza by airborne particles and to understand whether or not aerosols are likely to be an important route of transmission of influenza during a pandemic.

The purpose of this study is to gain a better understanding of the production and dissemination of aerosols containing the influenza virus. The results of this research will give scientists and health professionals greater insight into the airborne transmission of influenza and allow them to better assess the potential effectiveness of preventive measures.

The first part of this study will measure the quantity and size distribution of aerosol droplets produced by people with influenza when they cough. To accomplish this, volunteers with influenza-like illness will be asked to provide an oral swab for influenza testing, and then will cough into a spirometer. The aerosol produced by each person will be measured using commercially-available instrumentation. The oral swabs will be processed after the aerosol experiments are completed.

The second part of this study will determine the amount and size of airborne particles containing influenza virus that are present in a hospital emergency department during influenza season. Health care workers will be recruited to wear small aerosol collection devices as they go about their normal duties. The collected samples will then be analyzed for influenza virus. Adult patients in the emergency department with influenza-like illness will be asked to provide an oral swab to test for the flu virus in order to estimate the number of potential sources of viralladen airborne particles. There will be no costs to study participants. The total estimated annualized burden hours are 35.

TOTAL ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondent	Form	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Specific Aim 1: Volunteers with influenza				
Initial participants Qualified participants	Health questionnaire Consent form	42 40	1	5/60 20/60
Specific Aim 2: Health care workers				
Initial participants Qualified participants	Health questionnaire Consent form	32 30	1	5/60 20/60
Specific Aim 2: Emergency Department patients		•		
Participants	Consent form	15	1	20/60

Dated: December 14, 2007.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention. [FR Doc. E7–24929 Filed 12–21–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day-08-06BU]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Acting Reports Clearance Officer at 404–639–5960 and or send an e-mail to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395– 6974. Written comments should be received within 30 days of this notice.

Proposed Project

The Effectiveness of Teen Safe Driving Messages and Creative Elements on Parents and Teens—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Car crashes are the number one killer of teens, accounting for approximately one-third of all deaths within this age group. The National Center for Health Statistics reports that in 2004, a total of 3,620 young drivers were killed and an additional 303,000 were injured in motor vehicle crashes. In order to reduce these preventable deaths and injuries, parental awareness and education about Graduated Driver's Licensing (GDL) laws and the ways that parents can influence their children's safe driving are necessary. In preparation for a national campaign to educate parents about their role in their teens' driver education, it is necessary to determine the most effective messages and channels through which to communicate with parents. Ogilvy Public Relations Worldwide, PerformTech, International Communications Research (ICR) Survey and Fieldwork Network, on behalf of CDC, will conduct two studies to assess

the appropriateness and impact of messages and creative materials intended to (a) increase parental involvement in their teen's driving education and experience, and (b) encourage teens to adopt safer driving practices.

The first information collection will be accomplished through focus group testing of campaign messages and materials with representatives from our target audiences, parents and teens, in two cities in the U.S. The findings will provide valuable information regarding parents' and teens' levels of awareness and concern about safe driving; motivators for behavior change, especially GDL compliance; and message/channel preferences. The information collected will be used to develop final creative materials to implement the teen safe driving campaign in pilot cities. The second information collection will be accomplished through pilot city testing, which will evaluate knowledge, attitude and behaviors of intended audiences both pre- and post-communications campaign. The campaign will target parents of newly licensed drivers. It will encourage parents to understand state regulations regarding new drivers, talk with their teens about safe driving practices, and both manage and monitor their teens' driving behavior. Testing