#### Kimberly S. Lane,

Deputy Director, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2012–14396 Filed 6–12–12; 8:45 am] BILLING CODE 4163–18–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Disease Control and Prevention

[30-Day-12-12EG]

# 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) 639–7570 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) 395–5806. Written comments should be received within 30 days of this notice.

### Proposed Project

Use of Smartphones to Collect Information about Health Behaviors: Feasibility Study—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Despite the high level of public knowledge about the adverse effects of

smoking, tobacco use remains the leading preventable cause of disease and death in the U.S., resulting in approximately 443,000 deaths annually. During 2005–2010, the overall proportion of U.S. adults who were current smokers declined from 20.9% to 19.3%. Despite this decrease, smoking rates are still well above Healthy People 2010 targets for reducing adult smoking prevalence to 12%, and the decline in prevalence was not uniform across the population. Timely information on tobacco usage is needed for the design, implementation, and evaluation of public health programs.

The evolution of completely new, completely mobile communications technologies provides a unique opportunity for innovation in public health. Text messaging and smartphone web access are immediate, accessible, and anonymous, a combination of features that could make smartphones ideal for the ongoing research, surveillance, and evaluation of risk behaviors and health conditions, as well as targeted dissemination of information.

CDC proposes to conduct a feasibility study to identify and evaluate the process of conducting surveys by text message and smartphone, the outcomes of the surveys, and the value of the surveys. The universe for this study is English-speaking U.S. residents aged 18–65. The sample frame will consist of a national random digit dial sample of telephone numbers from a frame of known cell phone exchanges. Respondents reached on their cell phones will be asked to complete an initial CATI survey consisting of a short series of simple demographic questions, general health questions, and questions about tobacco and alcohol use. At the

conclusion of this brief survey, respondents who have smartphones will be asked to participate in the feasibility study, which consists of a first followup survey and, a week later, a second follow-up survey. Those who agree will receive invitations to participate by text message, which will include a link to the survey. A sample of respondents who do not have smartphones will be asked to participate in a text message pilot, which also consists of a first follow-up survey and a second followup survey. Text message respondents will receive a text message inviting them to participate; respondents opting in will be texted survey questions one at a time. Before initiating the feasibility study, CDC will conduct a brief pre-test of information collection forms and procedures.

This study will evaluate: (1) Response bias of a smartphone health survey by comparing data collected via CATI to data collected via smartphones/text messages, and data collected via smartphones to data collected via text messages, (2) relative cost-effectiveness of data collected via CATI to data collected via smartphones/text messages; (3) coverage bias associated with restricting the sample to smartphone users; and (4) the utility of smartphones for completing frequent, short interviews (e.g., diary studies to track activities or events).

OMB approval is requested for one year. Participation is voluntary and respondents can choose not to participate at any time. There are no costs to respondents other than their time. The total estimated annualized burden hours are 236.

## ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	No. of respondents	No. of responses per respondent	Avg. burden per response (in hr)
Adults Aged 18 to 65, All cell phone users	Pre-test (CATI Screener/CATI Recruitment	20	1	8/60
	CATI Screener	1,990	1	1/60
	CATI Recruitment	995	1	7/60
Adults Aged 18 to 65, Smartphone Users	First Web Survey Follow-up for Smartphone Users.	697	1	3/60
	Second Web Survey Follow-up for non- Smartphone Users.	592	1	3/60
Adults Aged 18 to 65, Non-smartphone Users	First Text Message Survey Follow-up for non-Smartphone Users.	200	1	3/60
	Second Text Message Survey Follow-up for non-Smartphone Users.	170	1	3/60

#### Kimberly S. Lane,

Deputy Director, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2012–14395 Filed 6–12–12; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Centers for Disease Control and Prevention

[60Day-12MX]

### 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 S. Lane, at 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

Research to Inform the Prevention of Asthma in Healthcare—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Healthcare is the largest industry in the United States and performs a vital function in society. Evidence from both surveillance and epidemiologic research

indicates that healthcare workers have an elevated risk for work-related asthma (WRA) associated with exposure to groups of agents such as cleaning products, latex, indoor air pollution, volatile organic compounds (VOCs) and bioaerosols. Recent epidemiologic studies of WRA among healthcare workers have utilized job exposure matrices (JEMs) based on probability of exposure, however, specific exposures/ etiologic agents are not well characterized and quantitative exposure measurements are lacking. In this project, NIOSH will augment the existing JEM with quantitative exposure data, which will significantly enhance the existing JEMs and develop a survey questionnaire for asthma in healthcare.

Since asthma continues to be a problem among healthcare workers, the overall goal of this project is to prevent work-related asthma among healthcare workers. The primary objective is to identify modifiable occupational risk factors for asthma in healthcare that will inform strategies for prevention. Specific Aims that support the Primary Objective are:

Aim 1. Measure frequency of asthma onset, related symptoms, and exacerbation of asthma in selected healthcare occupations.

Aim 2. Assess associations between asthma outcomes and exposures to identify modifiable risk factors.

In order to accomplish the goal and aims of this project, NIOSH has developed a survey designed to collect information about work history, workplace exposures and asthma health from workers in the healthcare industry. Aim 1 of this project will be completed using data exclusively from this survey. While aim 2 will be completed using asthma outcome data from the survey and exposure data from the JEM developed from survey data and exposure data from previously environmental sampling at healthcare facilities.

Approximately 17,500 health care workers in the New York City area will be recruited for this study. NIOSH is partnering with the Service Employees International Union (SEIU) Local 1199 in New York City. The SEIU1199 Communications Center (CC) will be responsible for collecting survey data from union members by telephone interview. The goal is to conduct a cross-sectional epidemiologic survey of approximately 5,000 healthcare workers who are members of SEIU1199. Only health care workers whose job titles are

in one of nine job titles will be recruited. These nine job titles include: certified nursing assistants (CNAs), central supply, environmental services, licensed practical nurses (LPNs), lab techs, operating room (OR) techs, registered nurses (RNs), respiratory therapists, and dental assistants. Furthermore, recruitment of health care workers will only be from hospitals and nursing homes.

Completion of the survey by SEIU1199 members will be done either online or over the telephone. After the initial recruitment period, SEIU1199 members will have approximately two weeks to complete the online survey. After this two week period, the SEIU1199 Communication Center will begin calling members who have not completed the online survey and attempt to complete the survey with them by telephone interview. NIOSH anticipates 20% of the responses to be made using the online survey and the remaining 80% to be by telephone interview.

Summary results of this study will be made available to SEIU1199 members who completed the survey through a letter mailed to their homes. Although NIOSH has partnered with SEIU119, results of this study will also be disseminated to other industry stakeholders including healthcare workers, researchers, clinicians, and professional societies and government agencies. The desired outcome of the dissemination efforts include healthcare workers learning about hazards in their work environment and becoming more prepared to participate in the development of strategies to minimize risk. Also, clinicians will learn how occupational exposures can impact the respiratory health of their patients who work in healthcare, which should improve the care they provide. In addition, manuscripts of results and conclusions will be drafted and published in peer reviewed journals.

The target sample size for this study is 5,000. Based on the SEIU1199 membership data, the percentage of eligible union members that fall into the targeted nine job categories is known. Therefore, a participant job-category distribution estimate can be made.

Completion of either the online or telephone survey will take approximately 30 minutes. It is estimated that the annualized burden will be 2,500 hours. There is no cost to respondents other than their time.