

Dated: April 20, 2016.

**Lorin S. Curit,**

Director, Federal Acquisition Policy Division,  
Office of Governmentwide Acquisition Policy,  
Office of Acquisition Policy, Office of  
Governmentwide Policy.

[FR Doc. 2016-09549 Filed 4-25-16; 8:45 am]

**BILLING CODE 6820-EP-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-16-16ACN; Docket No. CDC-2016-  
0038]

#### Proposed Data Collection Submitted for Public Comment and Recommendations

**AGENCY:** Centers for Disease Control and  
Prevention (CDC), Department of Health  
and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** The Centers for Disease  
Control and Prevention (CDC), as part of  
its continuing efforts to reduce public  
burden and maximize the utility of  
government information, invites the  
general public and other Federal  
agencies to take this opportunity to  
comment on proposed and/or  
continuing information collections, as  
required by the Paperwork Reduction  
Act of 1995. This notice invites  
comment on Workplace Health In  
America, a nationally representative  
survey of employer-based workplace  
health programs to describe the current  
state of U.S. workplace health  
promotion and protection programs and  
practices in employers of all sizes,  
industries and regions.

**DATES:** Written comments must be  
received on or before June 27, 2016.

**ADDRESSES:** You may submit comments,  
identified by Docket No. CDC-2016-  
0038 by any of the following methods:  
Federal eRulemaking Portal:  
Regulation.gov. Follow the instructions  
for submitting comments. Mail: Leroy A.  
Richardson, Information Collection  
Review Office, Centers for Disease  
Control and Prevention, 1600 Clifton  
Road NE., MS-D74, Atlanta, Georgia  
30329.

*Instructions:* All submissions received  
must include the agency name and  
Docket Number. All relevant comments  
received will be posted without change  
to Regulations.gov, including any  
personal information provided. For  
access to the docket to read background  
documents or comments received, go to  
Regulations.gov.

**Please note:** All public comment should be  
submitted through the Federal eRulemaking  
portal (Regulations.gov) or by U.S. mail to the  
address listed above.

**FOR FURTHER INFORMATION CONTACT:** To  
request more information on the  
proposed project or to obtain a copy of  
the information collection plan and  
instruments, contact the Information  
Collection Review Office, Centers for  
Disease Control and Prevention, 1600  
Clifton Road, NE., MS-D74, Atlanta,  
Georgia 30329; phone: 404-639-7570;  
Email: omb@cdc.gov.

**SUPPLEMENTARY INFORMATION:** Under the  
Paperwork Reduction Act of 1995 (PRA)  
(44 U.S.C. 3501-3520), Federal agencies  
must obtain approval from the Office of  
Management and Budget (OMB) for each  
collection of information they conduct  
or sponsor. In addition, the PRA also  
requires Federal agencies to provide a  
60-day notice in the **Federal Register**  
concerning each proposed collection of  
information, including each new  
proposed collection, each proposed  
extension of existing collection of  
information, and each reinstatement of  
previously approved information  
collection before submitting the  
collection to OMB for approval. To  
comply with this requirement, we are  
publishing this notice of a proposed  
data collection as described below.

*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; (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; and (e) estimates of capital  
or start-up costs and costs of operation,  
maintenance, and purchase of services  
to provide information. Burden means  
the total time, effort, or financial  
resources expended by persons to  
generate, maintain, retain, disclose or  
provide information to or for a Federal  
agency. This includes the time needed  
to review instructions; to develop,  
acquire, install and utilize technology  
and systems for the purpose of  
collecting, validating and verifying  
information, processing and  
maintaining information, and disclosing  
and providing information; to train  
personnel and to be able to respond to  
a collection of information, to search  
data sources, to complete and review

the collection of information; and to  
transmit or otherwise disclose the  
information.

#### Proposed Project

CDC Workplace Health Promotion  
Resource Center—New—National  
Center for Chronic Disease Prevention  
and Health Promotion (NCCDPHP),  
Centers for Disease Control and  
Prevention (CDC).

#### Background and Brief Description

The United States faces an  
unparalleled epidemic of poor health,  
driven largely by chronic diseases and  
conditions. A large body of literature  
shows that poor health, preceded by  
high levels of modifiable risk factors, is  
directly correlated with higher health  
care costs. Chronic conditions affect the  
workplace through health care costs,  
employee absences, safety claims, and  
presenteeism (*i.e.*, decrements in job  
performance due to health problems).

Workplaces are becoming important  
settings for health improvement and risk  
reduction. By improving the work  
health environment and helping  
workers achieve long-term behavior  
change, employers can diminish  
employees' risks for illnesses, enhance  
their quality of life, improve morale,  
eliminate unnecessary health care  
spending, minimize absences from  
work, reduce accidents, and increase  
productivity. Furthermore, having a  
healthy and productive workforce  
within a supportive work environment  
can foster greater loyalty among  
workers, a more committed workforce,  
and reduced turnover rates.

Despite their interest in improving the  
health and well-being of American  
workers, public and private employers  
often lack the know-how to do so  
effectively. A need exists for a trusted  
resource center housed in a virtual  
informational clearinghouse (IC) where  
employers and other stakeholders can  
access credible research (including best  
and promising practices), tools and  
resources, and technical assistance.

CDC plans to conduct information  
collection needed to design and  
implement a new CDC Workplace  
Health Promotion Resource Center  
(Resource Center), where relevant  
resources will be vetted, catalogued,  
compiled, and made publicly available  
to employers and other key  
stakeholders. Through the Resource  
Center, CDC will also provide technical  
assistance (TA) to employers, with the  
ultimate aim of improving population  
health, reducing health care utilization,  
and improving the productivity of  
employees. These activities are  
consistent with CDC's role as the

primary Federal agency for protecting health and promoting quality of life through the prevention and control of disease, injury, and disability. The CDC Workplace Health Promotion Resource Center is authorized by the Public Health Service Act and funded through the Prevention and Public Health Fund of the Patient Protection and Affordable Care Act (ACA).

Resource Center development and information collection will be conducted in two phases over a three-year period. In Phase 1 (project years 1 and 2), CDC will conduct formative research to understand the needs and preferences of the target audience. In Phase 2 (project years 2 and 3), CDC will build the Resource Center and IC, provide technical assistance, and assess customer satisfaction.

During Phase 1, CDC will conduct telephone interviews with 50 individuals who represent key Resource Center audiences: Employers (N=10), business groups (N=10), vendors and consultants (N=12), public health organizations (N=4), journalists (N=4), and researchers (N=10). Each tailored interview will be 45–60 minutes in length. Additional information will be collected through an online Needs and Interests Market Survey involving 800 respondents. Findings will be used to tailor the contents, technical support and dissemination practices of the Resource Center to the needs and interests of the target audiences.

During Phase 2, Resource Center products will be launched and CDC will collect brief, online customer satisfaction surveys from approximately

850 users. CDC will also pilot test and evaluate a direct technical assistance component of the Resource Center with approximately 5 selected states using two online surveys: a TA feedback survey and TA pilot assessment. The TA feedback survey will be offered to up to 100 stakeholders after each TA encounter and will take approximately 5 minutes. The TA pilot assessment will be provided at the conclusion of the TA pilot to up to 100 stakeholders and will take approximately 20 minutes. Findings will be used to improve workplace health programs and the offerings of the Resource Center.

OMB approval is requested for three years. Participation is voluntary and there are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden (in hrs.)
Employers .....	Needs and Interests Interview Guide for Employers.	3	1	1	3
Business Groups, Vendors, Consultants, and Public Health Organizations.	Needs and Interests Interview Guide for Business Groups, Vendors, Consultants, and Public Health Organizations.	9	1	1	9
Journalists .....	Needs and Interests Interview Guide for Journalists.	1	1	45/60	1
Researchers .....	Needs and Interests Interview Guide for the Research Community.	3	1	45/60	2
Key Stakeholders and Users of the Resource Center (All Groups).	Stakeholder Needs and Interests Market Survey.	267	1	20/60	89
Technical Assistance (TA) Participants .....	Consumer Satisfaction Survey .....	283	1	2/60	9
	TA Feedback Survey .....	33	5	5/60	14
	TA Pilot Assessment .....	33	1	20/60	11
<b>Total .....</b>					<b>138</b>

**Leroy A. Richardson,**  
 Chief, Information Collection Review Office,  
 Office of Scientific Integrity, Office of the  
 Associate Director for Science, Office of the  
 Director, Centers for Disease Control and  
 Prevention.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60Day-16-0199; Docket No. CDC-2016-0039]

**Proposed Data Collection Submitted for Public Comment and Recommendations**

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on an extension request for the information collection entitled *Application for Permit to Import Biological Agents and Vectors of Human Disease into the United States* and *Application for Permit to Import or Transport Live Bats (42 CFR 71.54)*.

**DATES:** Written comments must be received on or before June 27, 2016.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2016-0039 by any of the following methods:

- **Federal eRulemaking Portal:** Regulation.gov. Follow the instructions for submitting comments.

- **Mail:** Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road, NE., MS-D74, Atlanta, Georgia 30329.

**Instructions:** All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to Regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to Regulations.gov.

**Please note:** All public comment should be submitted through the Federal eRulemaking portal (Regulations.gov) or by U.S. mail to the address listed above.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of