

name, company name (if any), and “Information Collection 9000–0026, Change Order Accounting” on your attached document.

- **Mail:** General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0026, Change Order Accounting.

**Instructions:** Please submit comments only and cite Information Collection 9000–0026, Change Order Accounting, in all correspondence related to this collection. Comments received generally will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check [www.regulations.gov](http://www.regulations.gov), approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

**FOR FURTHER INFORMATION CONTACT:** Mr. Michael O. Jackson, Procurement Analyst, Office of Governmentwide Acquisition Policy, GSA, 202–208–4949, or email [michaelo.jackson@gsa.gov](mailto:michaelo.jackson@gsa.gov).

#### SUPPLEMENTARY INFORMATION:

##### A. Purpose

FAR 43.205 allows a contracting officer, whenever the estimated cost of a change or series of related changes under a contract exceeds \$100,000, to assert the right in the clause at FAR 52.243–6, Change Order Accounting, to require the contractor to maintain separate accounts for each change or series of related changes. Each account shall record all incurred segregable, direct costs (less allocable credits) of work, changed and unchanged, allocable to the change. These accounts are to be maintained until the parties agree to an equitable adjustment for the changes or until the matter is conclusively disposed of under the Disputes clause. This requirement is necessary in order to be able to account properly for costs associated with changes in supply and research and development contracts that are technically complex and incur numerous changes.

##### B. Annual Reporting Burden

*Respondents:* 8,850.  
*Responses per Respondent:* 12.  
*Annual Responses:* 106,200.  
*Hours per Response:* 1.  
*Total Burden Hours:* 106,200.

##### C. Public Comments

*Public comments are particularly invited on:* Whether this collection of

information is necessary for the proper performance of functions of the Federal Acquisition Regulations (FAR) and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

**Obtaining Copies Of Proposals:** Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755.

Please cite OMB Control No. 9000–0026, Change Order Accounting, in all correspondence.

Dated: January 15, 2016.

**Lorin S. Curit**

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

[FR Doc. 2016–01192 Filed 1–20–16; 8:45 am]

**BILLING CODE 6820–EP–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[60Day–16–16KA; Docket No. CDC–2016–0011]**

#### 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 a newly proposed information collection project entitled “Monitoring and Coordinating Personal Protective Equipment (PPE) in

Healthcare to Enhance Domestic Preparedness for Ebola Response”. The development of an ongoing Personal Protective Technology (PPT) sentinel surveillance system in the hospital setting will document data used to evaluate and monitor use and effectiveness for PPE usage in healthcare workers including Ebola protection.

**DATES:** Written comments must be received on or before March 21, 2016.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC–2016–0011 by any of the following methods:

Federal eRulemaking Portal: [Regulations.gov](http://www.regulations.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](http://www.regulations.gov), including any personal information provided. For access to the docket to read background documents or comments received, go to [Regulations.gov](http://www.regulations.gov).

**Please note:** All public comment should be submitted through the Federal eRulemaking portal ([Regulations.gov](http://www.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](mailto: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

Monitoring and Coordinating Personal Protective Equipment (PPE) in Healthcare to Enhance Domestic Preparedness for Ebola Response—New—National Center for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

### Background and Brief Description

The National Institute for Occupational Safety and Health (NIOSH) has the authority under the Occupational Safety and Health Act [29 CFR 671] to “develop recommendations for health and safety standards”, to “develop information on safe levels of exposure to toxic materials and harmful physical agents and substances”, and to “conduct research on new safety and health problems”. There is growing national concern for better understanding of the particular personal protective equipment (PPE) needs of healthcare workers to ensure the health and safety of this workforce during times of pandemic disease or bioterrorist threat. The use and effectiveness of the proper PPE are paramount to the management and mitigation of the effects of a disaster. NIOSH is requesting a three approval

from OMB to develop an ongoing PPT sentinel surveillance system in the hospital setting that will document data used to evaluate and monitor use and effectiveness for PPE usage in healthcare workers including Ebola protection.

NIOSH conducted a pilot study and partnered with four hospitals where respirator-related data were collected from a variety of stakeholders (less than 10 respondents) including Infection Control, Occupational Health, Emergency Preparedness, Environmental Health & Safety, and Purchasing. Surveillance metrics were established and shared with pilot participants on a regular basis throughout the pilot. Partners identified key performance indicators that this data might provide, such as the average number of respirators used per isolation order in the hospital, and identification of stakeholders and protocols impacting effective respirator use. Recommendations were made for monitoring schedules and survey improvement. The data collected during the pilot study provided experience and knowledge of respirator selection, availability, fit testing, usage patterns, outcomes, and confounders of respirator use and effectiveness at the four participating hospitals.

NIOSH now seeks approval to execute an approach for a minimum viable product (MVP) multi-hospital (15–20), real-time monitoring phase. The 15–20 facilities shall reflect the tiered approach recommended by CDC involving Frontline Healthcare Facilities, Ebola Assessment Hospitals and Ebola Treatment Centers. The effort shall be built upon the experience and knowledge obtained from the pilot projects, and shall be structured as the next step in the establishment of a national system to monitor usage and training for PPE used to protect against the Ebola virus based on current CDC recommendations. With this effort, the contractor shall develop and deploy the system to include a contingent of the domestic acute healthcare facilities in this three tier approach. The system content shall include status information for all PPE categories identified for protection against the hazards of Ebola exposure. The system will use a general interface engine designed to accept, validate, and process data from multiple, disparate sources.

The system will be developed to identify PPE replenishment needs to facilitate local, state, and eventually regional resource sharing and local purchasing as needed. It will also be compatible with PPE previously used at these facilities to allow seamless

continuity of patient care and worker protection. This capacity will offer a much-improved process for monitoring and maintaining appropriate PPE supplies through the constant, real-time monitoring of user demand, thus avoiding the misdirection of tens of millions of dollars' worth of respirators and other PPE to facilities that may not use distributed supplies due to a mismatch between products typically used and the supplies provided.

Respondents targeted for this study include hospital managers (also referred to in some cases as executives, coordinators or supervisors). These individuals are responsible for the day-to-day administration and/or implementation of the MVP. It is estimated that a sample of up to 20 hospitals will agree to participate among a variety of Ebola and Frontline treatment facilities. Participation will require no more than 255 minutes of workers' time per quarter. The hospitals will complete a baseline form and will also send quarterly and annual response as explained in the table below.

The Emergency and Crisis surveys are administered to hospitals via text message. The emergency survey is designed for an event spanning multiple weeks (e.g., pandemic). There are 3 preset questions that are related to Ebola and PPT supply concerns. The crisis survey is designed for an unanticipated scenario in which we may need to push ad hoc questions on a daily basis to hospitals. They will only be administered in a non-routine situation. During the 3 year approval period, we will test/train hospitals on each survey. However, they will not be part of the regular data collection.

### *Estimated Annualized Burden Hours*

The following is an explanation of the number of respondents for the annualized burden table. The baseline form is completed once by each hospital as they come onboard (20/3 = 7 rounded up). The annual form is completed by the hospitals in each year following their onboarding. Example: Year one, 5 hospitals onboarded; year two 6 new + 5 from previous year; year three 9 new + 11 from previous years. Thus, taking the sum of the previous year hospitals leads to 16 total (16/3 = 5 rounded down). The quarterly form is completed by all onboarded hospitals four times a year. The emergency and crisis forms are completed on all onboarded hospitals as needed but at least once for training and use the annualized number in the baseline form.

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden per respondent (hours)
Hospital .....	Baseline .....	7	1	8	56
Hospital .....	Annual .....	5	1	3	15
Hospital .....	Quarterly .....	12	4	3	144
Hospital .....	Emergency .....	7	4	15/60	7
Hospital .....	Crisis .....	7	7	10/60	8
Total .....	.....	.....	.....	.....	230

**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.

[FR Doc. 2016-01040 Filed 1-20-16; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-16-16KB; Docket No. CDC-2016-0010]

#### 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 a proposed information collection project entitled “Measuring Perceived Self-Escape Competencies among Underground Mineworkers”. The purpose of this two-year information collection is to gather survey data from up to 800 underground coal miners to measure their perceived competence in the critical knowledge, skills, and abilities that could be required for successful escape from an underground mine emergency.

**DATES:** Written comments must be received on or before March 21, 2016.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2016-0010 by any of the following methods:

Federal eRulemaking Portal: [Regulation.gov](http://www.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](http://www.Regulations.gov), including any personal information provided. For access to the docket to read background documents or comments received, go to [Regulations.gov](http://www.Regulations.gov).

**Please note:** All public comment should be submitted through the Federal eRulemaking portal ([Regulations.gov](http://www.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](mailto: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

Measuring Perceived Self-Escape Competencies among Underground Mineworkers—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety & health at work for all people through research and prevention. The Federal Mine Safety & Health Act of 1997, Public Law 91-173 as amended by Public Law 95-164, enables NIOSH to carry out research that is relevant to health and safety of workers in the underground coal industry. After a thorough review of United States' underground coal mine emergency escape preparedness and