be received within 60 days of this notice.

Proposed Project

Registration of Closed-Circuit Escape Respirator (CCER) units upon purchase—42 CFR part 84—Regulation—New—National Institute for Occupational Safety and Health (NIOSH), of the Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This project partially satisfies the requirement created by 42 CFR 84.311, Registration of CCER Units upon purchase. Applicants for approval of closed-circuit escape respirator (CCER) units must request respirator purchasers register their respirators with the National Institute for Occupational Safety and Health (NIOSH). The purpose of the information collection, is given in § 84.311c: ''The National Institute for Occupational Safety and Health (NIOSH) requests, but does not require, that purchasers of this respirator register each unit with NIOSH. Registration will enable NIOSH, which approved this model of respirator, to attempt to notify you if a problem is discovered that might affect the safety or performance of this respirator. Registration will also assist NIOSH in locating deployed units to periodically evaluate whether this respirator model is remaining effective under field conditions of storage and use.''

CCER units are respirators designed for escape from certain hazardous atmospheres, notably atmospheres that may be encountered during mining incidents. Subpart O, Closed-Circuit Escape Respirators, (§§ 84.300—84.311) was added to 42 CFR Part 84, Approval of Respiratory Protective Devices, describing requirements for a new class of NIOSH-approved respirators in response to issues with deployed Self-Contained Self-Rescuers (SCSR) respirators. Purchaser data collection was added to enable direct communication about potentially hazardous issues that may arise with approved CCER units, and to facilitate collection of CCER units from the field for evaluation.

In support of these goals, the collection will request the name and postal address of the company that purchased the respirators, a contact email address and position title, the respirator manufacturer, model, serial number or numbers, and date of manufacture, and the company industry and worksite regulation body (i.e. Mining Safety and Health Administration (MSHA), Occupational Safety and Health Administration (OSHA), or Other). Data collection will be through a structured email created using a NIOSH-hosted web form. Data collection is expected to take approximately five minutes per submission.

While the Federal Government is expected to purchase approximately 40,000 CCER units annually, these purchases will not be included in the burden estimate as MSHA will require the collection of this data for mine safety checks. Purchasers covered by MSHA regulations will be advised that MSHA reporting requirements will include all expected benefits of this CCER registration, and therefore registration is not recommended. The private sector is expected to purchase approximately 4,000 CCER units annually and a conservative estimate purchase lot size of ten (400 units).

We estimate an 80% response rate, for an estimated 320 responses. The estimated overall burden is 27 hours. There are no costs to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Responses per respondent	Average burden per response (in hours)	Total burden hours
Purchaser of CCER units	CCER Registration Form	320	1	5/60	27
Total					27

Leroy 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. 2014–16120 Filed 7–9–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–14–14AMW]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery—NEW—Centers for Disease Control and Prevention (CDC), National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP).

As part of a Federal Government-wide effort to streamline the process to seek feedback from the public on service delivery, the CDC has submitted a Generic Information Collection Request (Generic ICR): ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery'' to OMB for approval under the Paperwork Reduction Act (PRA) (44 U.S.C. 3501 et. seq.).

SUPPLEMENTARY INFORMATION:

Title: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

Abstract: The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training, or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative, and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior to fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

In accordance with 5 CFR 1320.8(d), Vol. 79, No. 83/Wednesday, April 30, 2014, a 60-day notice for public comment was published in the **Federal Register**. No public comments were received in response to this notice.

This is a new collection of information. Respondents will be screened and selected from Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government. Below we provide CDC's projected annualized estimate for the next three years. There is no cost to respondents other than their time. The estimated annualized burden hours for this data collection activity are 12,400.

ESTIMATED BURDEN HOURS

Type of collection	Number of respondents	Annual frequency per response	Hours per response	Total hours
Online, telephone surveys	15,000	1	30/60	7,500
Discussion groups	350	1	2	700
Focus groups	800	1	2	1,600
Website/app usability testing	2,000	1	30/60	1,000
Interviews	800	1	2	1,600

Leroy 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. 2014–16118 Filed 7–9–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Safety and Occupational Health Study Section: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Safety and Occupational Health Study Section, Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through June 30, 2016.

For more information contact: Price Connor, Ph.D., Executive Secretary, Safety and Occupational Health Study Section, Department of Health and Human Services, 1600 Clifton Road, NE., Mailstop E74, Atlanta, Georgia 30333, telephone 404/498–2511 or fax 404/498–2571.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Gary Johnson,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2014–16070 Filed 7–9–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Issuance and Enforcement Guidance for Dog Confinement Agreements

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

ACTION: Notice of Agency Guidance.