

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

OMB Number: To be determined.

Needs and Uses: The proposed information collection provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the agency's commitment to improving service delivery. Qualitative feedback means information that provides useful insights on perceptions and opinions, but is not a statistical survey that yields 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.

OGE expects to use various methods (e.g., focus groups, customer satisfaction surveys, comment cards), to solicit feedback. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public and other agency stakeholders. If this information is not collected, vital feedback from customers and stakeholders on the agency's services will be unavailable.

The agency will only submit a collection for approval under this generic clearance if it meets the following conditions:

- The collections are voluntary;
- The collections are low-burden for respondents (based on considerations of total burden hours, total number of respondents, or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government;
- The collections are non-controversial;
- The collections are focused on the awareness, understanding, attitudes, preferences, or experiences of the public or other stakeholders in order to improve existing or future services, products, or communication materials;
- Personally identifiable information (PII) is collected only to the extent necessary;
- Information gathered will be used only internally for general service improvement and program management purposes and is not intended for release to the public;

- Information gathered will not be used for the purpose of substantially informing influential policy decisions; and

- Information gathered will yield qualitative information; the collections will not be designed or expected to yield statistically reliable results or used as though the results are generalizable to the population of study.

Feedback collected under this generic clearance provides useful information, but it does 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.

As a general matter, information collections submitted under this generic clearance will not result in any new system of records containing privacy information and will not ask questions of a sensitive nature, such as sexual behavior and attitudes, religious beliefs, and other matters that are commonly considered private.

Current Action: New information collection request (generic).

Type of Review: New.

Affected Public: Individuals; Business or Other For-Profit Institutions; Not-For-Profit Institutions; State or Local Government.

Estimated Annual Number of Respondents: 45,000.

Projected average burden estimates for the next three years:

Average Expected Annual Number of Activities: 40.

Average Number of Respondents per Activity: 1,125.

Responses per Respondent: 1.

Annual Responses: 45,000.

Average Minutes per Response: 3 minutes.

Annual Burden Hours: 2,250 hours.

Frequency: On occasion.

Request for Comments: Agency and public comment is invited specifically on the need for and practical utility of this information collection, the accuracy of OGE's burden estimate, the enhancement of quality, utility and clarity of the information collected, and the minimization of burden (including the use of information technology). Comments received in response to this notice will be summarized for, and may be included with, the OGE generic information collection request. The comments will also become a matter of public record.

Approved: April 6, 2018.

David J. Apol,

General Counsel and Acting Director, U.S. Office of Government Ethics.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number CDC-2018-0033, NIOSH-311]

Draft—National Occupational Research Agenda for Public Safety

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Request for comment.

SUMMARY: The National Institute for Occupational Safety and Health of the Centers for Disease Control and Prevention announces the availability of a draft NORA Agenda entitled *National Occupational Research Agenda for Public Safety* for public comment. To view the notice and related materials, visit <https://www.regulations.gov> and enter CDC-2018-0033 in the search field and click "Search."

DATES: Electronic or written comments must be received by June 11, 2018.

ADDRESSES: You may submit comments, identified by CDC-2018-0033 and docket number NIOSH-311, by any of the following methods:

- *Federal eRulemaking Portal:* <https://www.regulations.gov> Follow the instructions for submitting comments.
 - *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C-34, Cincinnati, Ohio 45226-1998.
- Instructions:* All submissions received in response to this notice must include

the agency name and docket number [CDC–2018–0033; NIOSH–311]. All relevant comments received will be posted without change to <https://www.regulations.gov>, including any personal information provided. For access to the docket to read background documents or comments received, go to <https://www.regulations.gov>. All information received in response to this notice will also be available for public examination and copying at the NIOSH Docket Office, 1150 Tusculum Avenue, Room 155, Cincinnati, OH 45226–1998.

FOR FURTHER INFORMATION CONTACT:

Emily Novicki (*NORACoordinator@cdc.gov*), National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Mailstop E–20, 1600 Clifton Road NE, Atlanta, GA 30329, phone (404) 498–2581 (not a toll free number).

SUPPLEMENTARY INFORMATION: The National Occupational Research Agenda (NORA) is a partnership program created to stimulate innovative research and improved workplace practices. The national agenda is developed and implemented through the NORA sector and cross-sector councils. Each council develops and maintains an agenda for its sector or cross-sector.

Background: The National Occupational Research Agenda for Public Safety is intended to identify the research, information, and actions most urgently needed to prevent occupational injuries. The National Occupational Research Agenda for public safety provides a vehicle for stakeholders to describe the most relevant issues, gaps, and safety and health needs for the public safety sector. Each NORA research agenda is meant to guide or promote high priority research efforts on a national level, conducted by various entities, including: government, higher education, and the private sector.

The first National Occupational Research Agenda for Public Safety was published in 2009 for the second decade of NORA (2006–2016). This draft is an updated agenda for the third decade of NORA (2016–2026). The revised agenda was developed considering new information about injuries and illnesses, the state of the science, and the probability that new information and approaches will make a difference. As the steward of the NORA process, NIOSH invites comments on the draft *National Occupational Research Agenda for Public Safety*. Comments expressing support or with specific recommendations to improve the Agenda are requested. A copy of the draft Agenda is available at [https://](https://www.regulations.gov)

www.regulations.gov (see Docket Number CDC–2018–0033).

John J. Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–18–18UF; Docket No. CDC–2018–0032]

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 effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled *Evidence to Inform Standards that Ensure Turnout Gear Remains Protective Throughout Its Lifecycle* that will provide data that links turnout gear use conditions to its resulting performance characteristics.

DATES: CDC must receive written comments on or before June 11, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2018–0032 by any of the following methods:

- *Federal eRulemaking Portal:*

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. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all Federal comments 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 Leroy A. Richardson, 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 the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;
2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected; and
4. 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 submissions of responses.
5. Assess information collection costs.

Proposed Project

Evidence to Inform Standards that Ensure Turnout Gear Remains Protective Throughout Its Lifecycle—New—National Institute 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 been tasked to assure safe