DEPARTMENT OF DEFENSE
GENERAL SERVICES ADMINISTRATION
NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

Federal Acquisition Regulation; Submission for OMB Review; Contractors Performing Private Security Functions Outside the United States

AGENCIES: Department of Defense (DoD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice of request for an information collection requirement regarding a new OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve a new information collection requirement concerning Contractors Performing Private Security Functions Outside the United States. A notice was published in the Federal Register at 77 FR 43039, on July 23, 2012. No comments were received. 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.

DATES: Submit comments on or before December 27, 2012 to be considered in the formation of the final rule.

ADDRESS: Submit comments identified by Information Collection 9000–0184, Contractors Performing Private Security Functions Outside the United States, by any of the following methods:

- Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by inputting the OMB control number and selecting “Search”. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0184, Contractors Performing Private Security Functions Outside the United States”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0184, Contractors Performing Private Security Functions Outside the United States” on your attached document.
- Mail: General Services Administration, Regulatory Secretariat (MVCB), ATTN: Hada Flowers, 1275 First Street NE., 7th Floor, Washington, DC 20417.

Instructions: Please submit comments only and cite Information Collection 9000–0184, Contractors Performing Private Security Functions Outside the United States in all correspondence related to this case. All comments received will be posted without change to http://www.regulations.gov, including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Mr. Michael O. Jackson, Procurement Analyst, Governmentwide Acquisition Policy, at 202–206–4949 or email michaelo.jackson@gsa.gov.

SUPPLEMENTAL INFORMATION:

A. Purpose

Section 862 of the National Defense Authorization Act (NDAA) for Fiscal Year (FY) 2008, as amended by section 853 of the NDAA for FY 2009 and sections 831 and 832 of the NDAA for FY 2011, together with the required Governmentwide implementing regulations (32 CFR part 159, published at 76 FR 49650 on August 11, 2011), as amended, adds requirements and limitations for contractors performing private security functions in areas of contingency operations, combat operations, or other military operations as designated by the Secretary of Defense, upon agreement of the Secretaries of Defense and State. These requirements are that contractors performing in areas such as Iraq and Afghanistan ensure that their personnel performing private security functions comply with 32 CFR part 159, including (1) accounting for Government-acquired and contractor-furnished property and (2) reporting incidents in which a weapon is discharged, personnel are attacked or killed or property is destroyed, or active, lethal countermeasures are employed.

B. Annual Reporting Burden

Respondents: 920.

Responses per Respondent: 5.

Total Response: 4,600.

Hours per Response: 0.109 hours.

Total Burden Hours: 501.

Obtaining Copies of Proposals:

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street NE., Washington, DC 20417, telephone (202) 501–4753. Please cite OMB Control No. 9000–0184, Contractors Performing Private Security Functions Outside the United States, in all correspondence.

Dated: November 14, 2012.

William Clark,
Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

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/ATSDR Reports Clearance Office at (404) 639–7570 or send an email toomb@cdc.gov. Send written comments to CDC/ATSDR 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—Agency for Toxic Substances and Disease Registry (ATSDR).
As part of a Federal Government-wide effort to streamline the process to seek feedback from the public on service delivery, the ATSDR 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.).

To request additional information, please contact Kimberly S. Lane, Reports Clearance Officer, Centers for Disease Control and Prevention, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

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 qualitative 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 nonresponse 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.

The Agency received no comments in response to the 60-day notice published in the Federal Register on December 22, 2010 (75 FR 80542).

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 ATSDR’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 1,070.

<table>
<thead>
<tr>
<th>Type of collection</th>
<th>Average number of respondents per activity</th>
<th>Average frequency of response</th>
<th>Average number of activities</th>
<th>Average hours per response</th>
</tr>
</thead>
<tbody>
<tr>
<td>Comment cards or complaint forms</td>
<td>50</td>
<td>1</td>
<td>2</td>
<td>30/60</td>
</tr>
<tr>
<td>Focus groups</td>
<td>65</td>
<td>1</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>One-on-one interviews</td>
<td>50</td>
<td>1</td>
<td>1</td>
<td>30/60</td>
</tr>
<tr>
<td>One-time or panel discussion groups</td>
<td>10</td>
<td>1</td>
<td>2</td>
<td>8</td>
</tr>
<tr>
<td>Moderated, unmoderated, in-person and remote usability studies</td>
<td>500</td>
<td>1</td>
<td>1</td>
<td>30/60</td>
</tr>
<tr>
<td>Testing of a survey or other collection to refine questions</td>
<td>1,000</td>
<td>1</td>
<td>1</td>
<td>15/60</td>
</tr>
</tbody>
</table>

Dated: November 19, 2012.

Ron A. Otten,
Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2012–28741 Filed 11–26–12; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–13–0914]

Agency Forms Undergoing Paperwork Reduction Act Review

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Proposed Project

Workplace Violence Prevention Programs in NJ Healthcare Facilities (0920–0914, Expiration 1/31/2015)—Revision—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Healthcare workers are nearly five times more likely to be victims of violence than workers in all industries combined. While healthcare workers are not at particularly high risk for job-related homicide, nearly 60% of all nonfatal assaults occurring in private industry are experienced in healthcare.

Six states have enacted laws to reduce violence against healthcare workers by requiring workplace violence prevention programs. However, little is understood about how effective these laws are in reducing violence against healthcare workers.

The objective of the proposed study is three-fold: (1) To examine healthcare facility compliance with the New Jersey Violence Prevention in Health Care Facilities Act, (2) to evaluate the effectiveness of the regulations in this Act in reducing assault injuries to workers. Our central hypothesis is that facilities with high compliance with the regulations will have lower rates of