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. 9000–0101, Drug-Free Workplace, in all correspondence.

Edward Loeb,
Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2015–31561 Filed 12–15–15; 8:45 am]
BILLING CODE 6820–EP–P

DEPARTMENT OF DEFENSE
GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0083; Docket 2015–0055; Sequence 31]

Information Collection; Qualification Requirements

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

ACTION: Notice of request for comments regarding the extension of a previously existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning Qualification Requirements.

DATES: Submit comments on or before February 16, 2016.

ADDRESSES: Submit comments identified by Information Collection 9000–0083, Qualification Requirements, by any of the following methods:
- Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0083, Qualification Requirements”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0083, Qualification Requirements” on your attached document.
- Mail: General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, ATTN: Ms. Flowers/IC 9000–0083, Qualification Requirements.

Instructions: Please submit comments only and cite Information Collection 9000–0083, Qualification Requirements, 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, 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. Charles Gray, Procurement Analyst, Office of Governmentwide Acquisition Policy, GSA, 703–795–6328 or charles.gray@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

FAR subpart 9.2 and the associated clause at FAR 52.209–1, implement the statutory requirements of 10 U.S.C. 2319 and 41 U.S.C. 3311, which allow an agency to establish a qualification requirement for testing or other quality assurance demonstration that must be completed by an offeror before award of a contract. Under the qualification requirements, an end item, or a component thereof, may be required to be prequalified.

The clause at FAR 52.209–1, Qualification Requirements, requires offerors who have met the qualification requirements to identify the offeror’s name, the manufacturer’s name, source’s name, the item name, service identification, and test number (to the extent known). This eliminates the need for an offeror to provide new information when the offeror, manufacturer, source, product or service covered by qualification requirement has already met the standards specified by an agency in a solicitation.

The contracting officer uses the information to determine eligibility for award when the clause at 52.209–1 is included in the solicitation. Alternatively, items not yet listed may be considered for award upon the submission of evidence of qualification with the offer.

B. Annual Reporting Burden

Respondents: 9,693.

Responses per Respondent: 5.


C. Public Comments

Public comments are particularly invited on: Whether the collection of information is necessary; 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. 9000–0083, Qualification Requirements, in all correspondences.

Edward Loeb,
Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2015–31560 Filed 12–15–15; 8:45 am]
BILLING CODE 6820–EP–P

DEPARTMENT OF DEFENSE
GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[Docket 2015–0055; Sequence 16; OMB Control No. 9000–0107]

Submission for OMB Review; Notice of Radioactive Materials

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

ACTION: Notice of request for comments regarding the extension of a previously existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning Notice of Radioactive Materials. A
notice was published in the Federal Register at 80 FR 58253 on September 28, 2015. No comments were received.

DATES: Submit comments on or before January 13, 2016.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: Office of Information and Regulatory Affairs of OMB, Attention: Desk Officer for GSA, Room 10236, NEOB, Washington, DC 20503. Additionally submit a copy to GSA by any of the following methods:

- **Regulations.gov**: Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0107, Notice of Radioactive Materials”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0107, Notice of Radioactive Materials” on your attached document.
- **Mail**: General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0107, Notice of Radioactive Materials.

Instructions: Please submit comments only and cite Information Collection 9000–0107, Notice of Radioactive Materials, 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, 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. Charles Gray, Procurement Analyst, Office of Acquisition Policy, GSA, 703–795–6328 or email charles.gray@gsa.gov.

A. Purpose

The clause at FAR 52.223–7, Notice of Radioactive Materials, requires contractors to notify the Government prior to delivery of items containing radioactive materials. The purpose of the notification is to alert receiving activities that appropriate safeguards may need to be instituted. The notice shall specify the part or parts of the items which contain radioactive materials, a description of the materials, the name and activity of the isotope, the manufacturer of the materials, and any other information known to the contractor which will put users of the items on notice as to the hazards involved.

B. Annual Reporting Burden

<table>
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<th>Respondents</th>
<th>Responses per Respondent</th>
<th>Annual Responses</th>
<th>Hours per Response</th>
<th>Total Burden Hours</th>
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<td>2,675</td>
<td>1</td>
<td>2,675</td>
</tr>
</tbody>
</table>

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FAR, and whether it will have practical utility; whether our estimate of the 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.


Edward Loeb,
Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2015–31562 Filed 12–15–15; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

Centers for Disease Control and Prevention

[30Day–16–0048]

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. Direct written comments and/or suggestions regarding the items contained in this notice 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.

Projected Project

ATSDR Exposure Investigations (EIs) (OMB Control No. 0923–0048, Expiration Date 5/31/2016)—Extension—Agency for Toxic Substances and Disease Registry (ATSDR).

Background and Brief Description

The Agency for Toxic Substances and Disease Registry (ATSDR) is requesting a three-year extension of this generic clearance to allow the agency to conduct exposure investigations (EIs), through methods developed by ATSDR. After a chemical release or suspected release into the environment, EIs are usually requested by officials of a state health agency, county health departments, the Environmental Protection Agency (EPA), the general public, and ATSDR staff.