DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0157; Docket 2014–0055; Sequence 17]

Federal Acquisition Regulation; Information Collection Architect-Engineer Qualifications (SF 330)

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

ACTION: Notice of request for public comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act (44 U.S.C. chapter 35), the Regulatory Secretariat Division (MVCB) 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 for the Architect–Engineer Qualifications form (SF 330).

DATES: Submit comments on or before June 6, 2014.

ADDRESSES: Submit comments identified by Information Collection 9000–0157 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 9000–0157. Select the link ''Comment Now'' that corresponds with ''Information Collection 9000–0157''. Follow the instructions provided on the screen. Please include your name, company name (if any), and ''Information Collection 9000–0157'' on your attached document.
- *Fax:* 202–501–4067.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0157.

Instructions: Please submit comments only and cite Information Collection 9000–0157, in all correspondence related to this collection. 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. Curtis E. Glover, Sr., Procurement Analyst, Contract Policy Division, GSA 202–501–1448 or email *Curtis.glover@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

Standard Form 330, Part I is used by all Executive agencies to obtain information from architect-engineer firms interested in a particular project. The information on the form is reviewed by a selection panel composed of professional people and assists the panel in selecting the most qualified architect-engineer firm to perform the specific project. The form is designed to provide a uniform method for architect-engineer firms to submit information on experience, personnel, capabilities of the architect-engineer firm to perform along with information on the consultants they expect to collaborate with on the specific project.

Standard Form 330, Part II is used by all Executive agencies to obtain general uniform information about a firm's experience in architect-engineering projects. Architect-engineer firms are encouraged to update the form annually. The information obtained on this form is used to determine if a firm should be solicited for architect-engineer projects.

B. Annual Reporting Burden

Respondents: 5,000.

Responses per Respondent: 4.

Total Responses: 20,000.

Hours per Response: 29.

Total Burden Hours: 580,000.

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 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. Please cite OMB Control No. 9000–0157, Architect-Engineer Qualifications (SF 330), in all correspondence.

Dated: April 1, 2014.

Karlos Morgan,

Acting Director, Federal Acquisition Policy Division, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2014–07625 Filed 4–4–14; 8:45 am]

BILLING CODE 6820–EP–P

GENERAL SERVICES ADMINISTRATION

[Notice–PMAB–2014–01; Docket No. 2014–0002; Sequence No.14]

The President's Management Advisory Board (PMAB); Notification of Upcoming Public Advisory Meeting

AGENCY: Office of Executive Councils, U.S. General Services Administration (GSA).

ACTION: Meeting notice.

SUMMARY: The President's Management Advisory Board (PMAB), a Federal Advisory Committee established in accordance with the Federal Advisory Committee Act (FACA), 5 U.S.C., App., and Executive Order 13538, will hold a public meeting on Friday, April 25, 2014.

DATES: *Effective:* April 7, 2014.

Meeting date: The meeting will be held on Friday, April 25, 2014 beginning at 9:00 a.m. eastern time, ending no later than 1:00 p.m.

ADDRESSES: Eisenhower Executive Office Building, 1650 Pennsylvania Avenue NW., Washington, DC.

FOR FURTHER INFORMATION CONTACT: Mr. Stephen Brockelman, Designated Federal Officer, President's Management Advisory Board, Office of Executive Councils, General Services Administration, 1800 F Street NW., Washington, DC 20006, at *stephen.brockelman@gsa.gov.*

SUPPLEMENTARY INFORMATION:

Background: The PMAB was established to provide independent advice and recommendations to the President and the President's Management Council on a wide range of issues related to the development of effective strategies for the implementation of best business practices to improve Federal Government management and operation.

Agenda: The main purpose of this meeting is for the PMAB to discuss and define their focus area for 2014, improving customer service in the Federal government. The meeting will aim to identify the challenges and opportunities for customer service improvement in various Federal

agencies. The PMAB will also receive progress updates on management initiatives for which they issued recommendations in prior years. Finally, the meeting will cover planning and logistics for PMAB during the coming year.

Meeting Access: The PMAB will convene its meeting in the Eisenhower Executive Office Building, 1650 Pennsylvania Avenue NW., Washington, DC. Due to security, there will be no public admittance to the Eisenhower Building to attend the meeting. However, the meeting is open to the public; interested members of the public may view the PMAB's discussion at *http://www.whitehouse.gov/live.* Members of the public wishing to comment on the discussion or topics outlined in the Agenda should follow the steps detailed in Procedures for Providing Public Comments below.

Availability of Materials for the Meeting: Please see the PMAB Web site (*http://www.whitehouse.gov/administration/advisory-boards/pmab*) for any materials available in advance of the meeting and for meeting minutes that will be made available after the meeting. Detailed meeting minutes will be posted within 90 days of the meeting.

Procedures for Providing Public Comments: In general, public statements will be posted on the PMAB Web site (*http://www.whitehouse.gov/administration/advisory-boards/pmab*). Non-electronic documents will be made available for public inspection and copying in PMAB offices at GSA, 1800 F Street NW., Washington, DC 20006, on official business days between the hours of 10 a.m. and 5 p.m. eastern time. You can make an appointment to inspect statements by telephoning 202–501–1398. All statements, including attachments and other supporting materials, received are part of the public record and subject to public disclosure. Any statements submitted in connection with the PMAB meeting will be made available to the public under the provisions of the Federal Advisory Committee Act.

The public is invited to submit written statements for this meeting until 12:30 p.m. eastern time on Thursday, April 24, by either of the following methods: *Electronic or Paper Statements:* Submit electronic statements to Mr. Brockelman, Designated Federal Officer at *stephen.brockelman@gsa.gov;* or send paper statements in triplicate to Mr. Brockelman at the PMAB GSA address above.

Dated: April 1, 2014.

Anne Rung,

Associate Administrator, Office of Government-wide Policy, General Services Administration.

[FR Doc. 2014–07755 Filed 4–4–14; 8:45 am]

BILLING CODE 6820–BR–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–14–0109]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–7570 or send comments to LeRoy Richardson, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an email to *omb@cdc.gov.*

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; and (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. Written comments should be received within 60 days of this notice.

Proposed Project

Respiratory Protective Devices—42 CFR part 84—Regulation—(0920–0109)—Revision—National Institute for Occupational Safety and Health (NIOSH), of the Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This data collection was formerly named Respiratory Protective Devices 30 CFR part 11 but in 1995, the respirator standard was moved to 42 CFR Part 84. The regulatory authority for the National Institute for Occupational Safety and Health (NIOSH) certification program for respiratory protective devices is found in the Mine Safety and Health Amendments Act of 1977 (30 U.S.C. 577a, 651 et seq., and 657(g)) and the Occupational Safety and Health Act of 1970 (30 U.S.C. 3, 5, 7, 811, 842(h), 844). These regulations have, as their basis, the performance tests and criteria for approval of respirators used by millions of American construction workers, miners, painters, asbestos removal workers, fabric mill workers, and fire fighters.

Regulations of the Environmental Protection Agency (EPA) and the Nuclear Regulatory Commission (NRC) also require the use of NIOSH-approved respirators. These regulations also establish methods for respirator manufacturers to submit respirators for testing under the regulation and have them certified as NIOSH-approved if they meet the criteria given in the above regulation.

NIOSH, in accordance with 42 CFR Part 84: (1) Issues certificates of approval for respirators which have met specified construction, performance, and protection requirements; (2) establishes procedures and requirements to be met in filing applications for approval; (3) specifies minimum requirements and methods to be employed by NIOSH and by applicants in conducting inspections, examinations, and tests to determine effectiveness of respirators; (4) establishes a schedule of fees to be charged applicants for testing and certification, and (5) establishes approval labeling requirements. Information is collected from those who request services under 42 CFR Part 84 in order to properly establish the scope and intent of request.

Information collected from requests for respirator approval functions includes contact information and information about factors likely to affect respirator performance and use. Such information includes, but is not necessarily limited to, respirator design, manufacturing methods and materials, quality assurance plans and procedures, and user instruction and draft labels, as specified in the regulation.

The main instrument for data collection for respirator approval functions is the Standard Application for the Approval of Respirators (SAF), currently Version 7. A replacement instrument which will collect the same information is in development.

Respirator manufacturers are the respondents (estimated to average 63 each year over the years 2014–2016) and upon completion of the SAF their