If you file your comment on paper, write ''Subpart N of Regulation V, PRA Comment, P125403,'' on your comment and on the envelope. You can mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW., Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW., 5th Floor, Suite 5610 (Annex J), Washington, DC 20024.

The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before November 27, 2015. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed to Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW., Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead should be sent by facsimile to (202) 395–5806.

David C. Shonka,

Principal Deputy General Counsel.

[FR Doc. 2015–27446 Filed 10–27–15; 8:45 am]

BILLING CODE 6750–01–P

GENERAL SERVICES ADMINISTRATION

[Notice–PMAB–2015–02; Docket No. 2015–0002; Sequence 29]

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

AGENCY: Office of Executive Councils, 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), will hold a public meeting on Monday, November 16, 2015.

DATES: *Effective:* October 28, 2015.

Meeting date: The meeting will be held on Monday, November 16, 2015, beginning at 9:00 a.m. Eastern Standard Time (EST), ending no later than 1:00 p.m., EST.

ADDRESSES: The meeting will be held at the Eisenhower Executive Office Building, 1650 Pennsylvania Avenue NW., Washington, DC.

FOR FURTHER INFORMATION CONTACT: Mr. Brad Golson, Designated Federal Officer, President's Management Advisory Board, Office of Executive Councils, GSA, 1800 F Street NW., Washington, DC 20006, at 202–969–7989, or via email at *brad.golson@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 to obtain recommendations from PMAB members on effective implementation of the FedStat process used by the Office of Management and Budget (OMB), to assess effective management practices, and of the Federal Information Technology Acquisition Reform Act (FITARA), which was passed by Congress during the 113th session of the United Sates Congress.

Meeting Access

The PMAB will convene its meeting in the Eisenhower Executive Office Building, 1650 Pennsylvania Avenue NW., Washington, DC 20504. 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 and may be viewed 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 20405, on official business days between the hours of 10 a.m., and 5 p.m., EST. You can make an appointment to inspect statements by telephoning 202–695–9554. 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., EST, on Friday, November 13, by either of the following methods: *Electronic or Paper Statements:* Submit electronic statements to Mr. Golson, Designated Federal Officer at *brad.golson@gsa.gov;* or send paper statements in triplicate to Mr. Golson at the PMAB GSA address above.

Dated: October 21, 2015.

Christine Harada,

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

[FR Doc. 2015–27368 Filed 10–27–15; 8:45 am]

BILLING CODE 6820–BR–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–00XX; Docket No. 2015–0055; Sequence 49]

Information Collection; Payment to Small Business Subcontractors

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

ACTION: Withdrawal of notice.

SUMMARY: The notice, OMB Control No. 9000–00XX, Payment to Small Business

Subcontractors, published in the **Federal Register**, is being withdrawn and is no longer accepting comments.

DATES: *Effective:* October 28, 2015.

FOR FURTHER INFORMATION CONTACT: Mr. Curtis E. Glover, Sr., Procurement Analyst, GSA, at 202–501–1448, or via email to *curtis.glover@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The notice, published in the **Federal Register** at 80 FR 60383, on October 6, 2015, requesting comments regarding a new information collection, 9000–00XX; Payment to Small Business Subcontractors, is being withdrawn. The notice is being withdrawn because it is associated with a rule which is still in process, and has not been published. Comments are no longer being sought at this time; however, the public will have a chance to comment once the rule is published.

Edward Loeb,

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

[FR Doc. 2015–27432 Filed 10–27–15; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–0840]

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.

Proposed Project

Formative Research and Tool Development (OMB Control No. 0920–0840, Expiration 02/29/2016)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD, TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP) requests a three-year approval and extension of the ''Formative Research and Tool Development'' generic information collection plan. This information collection request is designed to allow NCHHSTP to conduct formative research information collection activities used to inform many aspects of surveillance, communications, health promotion, and research project development for NCHHSTP's 4 priority diseases (HIV/AIDS, sexually transmitted diseases/infections (STD/STI), viral hepatitis, tuberculosis elimination and the Division of School and Adolescent Heath (DASH).

Formative research is the basis for developing effective strategies including communication channels, for influencing behavior change. It helps researchers identify and understand the characteristics—interests, behaviors and needs—of target populations that influence their decisions and actions.

Formative research is integral in developing programs as well as improving existing and ongoing programs. Formative research also looks at the community in which a public health intervention is being or will be implemented and helps the project staff understand the interests, attributes and needs of different populations and persons in that community. Formative research is research that occurs before a program is designed and implemented, or while a program is being conducted.

NCHHSTP formative research is necessary for developing new programs or adapting programs that deal with the complexity of behaviors, social context, cultural identities, and health care that underlie the epidemiology of HIV/AIDS, viral hepatitis, STDs, and TB in the U.S, as well as for school and adolescent health.

CDC conducts formative research to develop public-sensitive communication messages and user friendly tools prior to developing or recommending interventions, or care. Sometimes these studies are entirely behavioral but most often they are cycles of interviews and focus groups designed to inform the development of a product.

Products from these formative research studies will be used for prevention of HIV/AIDS, Sexually Transmitted Infections (STI), viral Hepatitis, and Tuberculosis. Findings from these studies may also be presented as evidence to disease-specific National Advisory Committees, to support revisions to recommended prevention and intervention methods, as well as new recommendations.

Much of CDC's health communication takes place within campaigns that have fairly lengthy planning periods—timeframes that accommodate the standard Federal process for approving data collections. Short term qualitative interviewing and cognitive research techniques have previously proven invaluable in the development of scientifically valid and population-appropriate methods, interventions, and instruments.

This request includes studies investigating the utility and acceptability of proposed sampling and recruitment methods, intervention contents and delivery, questionnaire domains, individual questions, and interactions with project staff or electronic data collection equipment. These activities will also provide information about how respondents answer questions and ways in which question response bias and error can be reduced.

This request also includes collection of information from public health programs to assess needs related to initiation of a new program activity or expansion or changes in scope or implementation of existing program