This site is accessible to individuals with disabilities. The meeting also will be made available via teleconference and/or webinar.

Submit comments identified by ''Notice–CECANF–2015–01,'' by either of the following methods:

• Regulations.gov: *http://www.regulations.gov*.

Submit comments via the Federal eRulemaking portal by searching for ''Notice–CECANF–2015–01.'' Select the link ''Comment Now'' that corresponds with ''Notice–CECANF–2015–01.'' Follow the instructions provided on the screen. Please include your name, organization name (if any), and ''Notice–CECANF–2015–01'' on your attached document.

• Mail: U.S. General Services Administration, 1800 F Street NW., Room 7003D, Washington DC 20405, Attention: Tom Hodnett (CD) for CECANF.

Instructions: Please submit comments only and cite ''Notice–CECANF–2015–01'' in all correspondence related to this notice. 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: Visit the CECANF Web site at *https://eliminatechildabusefatalities.sites.usa.gov/* or contact Patricia Brincefield, Communications Director, at 202–818–9596, U.S. General Services Administration, 1800 F Street NW., Room 7003D, Washington DC 20405, Attention: Tom Hodnett (CD) for CECANF.

SUPPLEMENTARY INFORMATION:

Background: CECANF was established to develop a national strategy and recommendations for reducing fatalities resulting from child abuse and neglect.

Agenda: On February 26, 2015, Commission members will meet to hear testimony from four panels of presenters. The first panel will focus on service delivery models under way in the Pacific Northwest region that are promising in the prevention of fatalities to children *not known* to the child welfare system. During the second panel, Commission members will hear testimony on workforce issues that influence performance and successful strategies for addressing these issues. JooYeun Chang, Associate Commissioner of the Children's Bureau, will talk with Commissioners about the status and effectiveness of key federal policies aimed at protecting children at risk of harm. She will provide critical information regarding successful strategies that are being used by states, tribes, and localities that demonstrate reduced child abuse and neglect fatalities to children known to child welfare. In the fourth panel, Commissioners will learn about the legal framework that supports or limits government's ability to intervene on behalf of children in the family context. Commission members will then continue discussing the work plans of the six Commission subcommittees, the information that they have obtained to date, and emerging high-level recommendations.

Attendance at the Meeting: Individuals interested in attending the meeting in person or participating by webinar and teleconference must register in advance. To register to attend in person or by webinar/phone, please go to *https://attendee.gotowebinar.com/rt/842048149990446850* and follow the prompts. Once you register, you will receive a confirmation email with the webinar login and teleconference number. Detailed meeting minutes will be posted within 90 days of the meeting. Members of the public will not have the opportunity to ask questions or otherwise participate in the meeting.

However, members of the public wishing to comment should follow the steps detailed under the heading **ADDRESSES** in this publication or contact us via the CECANF Web site at *https://eliminatechildabusefatalities.sites.usa.gov/contact-us/*.

Dated: January 21, 2015.

Karen White,

Executive Assistant.

[FR Doc. 2015–02052 Filed 2–2–15; 8:45 am]

BILLING CODE 6820–34–P

GOVERNMENT ACCOUNTABILITY OFFICE

Health Information Technology Policy Committee Nomination Letters

AGENCY: Government Accountability Office (GAO).

ACTION: Notice on letters of nomination of candidates.

SUMMARY: The American Recovery and Reinvestment Act of 2009 (ARRA) established the Health Information Technology Policy Committee (Health IT Policy Committee) and gave the Comptroller General responsibility for appointing 13 of its 20 members. As the result of terms ending in April 2015, GAO is accepting nominations of individuals for four openings on the committee in the following categories of representation or expertise required in ARRA: Advocate for patients or consumers, health care provider, representative of a health plan or third party payer, and expertise in health care quality measurement and reporting. For appointments to the HIT Policy committee to be made in April 2015 in these categories, I am announcing the following: Letters of nomination and resumes should be submitted by February 27, 2015 to ensure adequate opportunity for review and consideration of nominees.

ADDRESSES: *Email: HITCommittee@gao.gov.*

Mail: ATTN: HITPC Appointments, U.S. GAO, 441 G Street NW., Washington, DC 20548.

FOR FURTHER INFORMATION CONTACT: GAO Office of Public Affairs, (202) 512–4800. 42 U.S.C. 300jj-12.

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2015–01837 Filed 2–2–15; 8:45 am]

BILLING CODE 1610–02–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the National Advisory Committee on Children and Disasters

AGENCY: Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) is hereby giving notice that the National Advisory Committee on Children and Disasters (NACCD) will be holding a meeting via teleconference. The meeting is open to the public.

DATES: The February 26, 2015, NACCD meeting is scheduled from 2:00 p.m. to 3:00 p.m. EST. The agenda is subject to change as priorities dictate. Please check the NACCD Web site, located at *WWW.PHE.GOV/NACCD* for the most up-to-date information on the meeting.

ADDRESSES: To attend the meeting via teleconference, call toll-free: 1–877–601–4720. The pass-code is: 9977742. Please call 15 minutes prior to the beginning of the conference call to facilitate attendance. Pre-registration is required for public attendance. Individuals who wish to attend the meeting should submit an inquiry via the NACCD Contact Form located at *www.phe.gov/NACCDComments.*

FOR FURTHER INFORMATION CONTACT: Please submit an inquiry via the NACCD Contact Form located at *www.phe.gov/NACCDComments.*

SUPPLEMENTARY INFORMATION: Pursuant to the Federal Advisory Committee Act (FACA) of 1972 (5 U.S. C., Appendix, as amended), and section 2811A of the Public Health Service (PHS) Act (42 U.S. C. 300hh–10a), as added by section 103 of the Pandemic and All Hazards Preparedness Reauthorization Act of 2013 (Pub. L. 113–5), the HHS Secretary, in consultation with the Secretary of the U.S. Department of Homeland Security, established the National Advisory Committee on Children and Disasters (NACCD). The purpose of the NACCD is to provide advice and consultation to the HHS Secretary with respect to the medical and public health needs of children in relation to disasters. The Office of the Assistant Secretary for Preparedness and Response (ASPR) provides management and administrative oversight to support the activities of the NACCD.

Background: This public meeting will be dedicated to the members voting to approve the report of findings of the NACCD Surge Capacity Work Group.

Availability of Materials: The meeting agenda and materials will be posted on the NACCD Web site at: *www.phe.gov/naccd* prior to the meeting.

Procedures for Providing Public Input: All written comments must be received prior to February 24, 2015. Please submit comments via the NACCD Contact Form located at *www.phe.gov/NACCDComments.* Individuals who plan to attend and need special assistance should submit a request via the NACCD Contact Form located at *www.phe.gov/NACCDComments.*

Dated: January 22, 2015.

Nicole Lurie,

Assistant Secretary for Preparedness and Response.

[FR Doc. 2015–01615 Filed 2–2–15; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–15–0940]

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*. Written comments and/or suggestions regarding the items contained in this notice should be directed 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

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB No. 0920–0940, exp. 06/30/2015)—Extension—Centers for Disease Control and Prevention (CDC), National Institute for Occupational Safety and Health (NIOSH), Health Hazard Evaluation Program.

As part of a Federal Government-wide effort to streamline the process to seek feedback from the public on service delivery, the CDC 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 Leroy A. Richardson, 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 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.

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 non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior 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 April 30, 2014 (75 FR 24432).

This is a new collection of information. Respondents will be