ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average hours per response (in hours)
Facility office staff approaching sampled persons for enrollment.	N/A	1,090	1	5/60
Facility office staff pulling medical records	N/A	8,720	1	3/60

Leroy A. Richardson

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2015–07839 Filed 4–6–15; 8:45 am]

BILLING CODE CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–15–15UK]

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—NEW—National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention (CDC).

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*.).

Background and Brief Description

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 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.

In accordance with 5 CFR 1320.8(d), Vol. 79, No. 83/Wednesday, April 30, 2014, a 60 day notice for public comment was published in the **Federal Register**. No public comments were received in response to this notice.

This is a new collection of information. Respondents will take online surveys or participate in Web site usability testing, interviews, discussion groups, or focus groups. Below is Centers for Disease Control and Prevention (CDC), National Center for Emerging and Zoonotic Infectious Diseases (NCEZID) projected 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 is 3,850 hours:

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Avgerage burden per response
Customer or Stakeholder	Online surveys	2,500	1	30/60
Customer or Stakeholder	Discussion Groups	150	1	120/60
Customer or Stakeholder	Focus groups	700	1	120/60
Customer or Stakeholder	Website/app usability testing	250	1	45/60
Customer or Stakeholder	Interviews	100	1	90/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2015–07840 Filed 4–6–15; 8:45 am]

BILLING CODE CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Health Resources and Services Administration

Advisory Committee on the Maternal, Infant and Early Childhood Home Visiting Program Evaluation

AGENCY: Administration for Children and Families (ACF), HHS; Health Resources and Services Administration (HRSA), HHS.

ACTION: Notice to announce the renewal of the Advisory Committee on the Maternal, Infant and Early Childhood Home Visiting Program Evaluation.

Authority: Sec. 511(g)(1) of Title V of the Social Security Act (42 U.S.C. 711, *et seq.*). The Committee is governed by provisions of Public Law 92–463, as amended, (5 U.S.C. App. 2), which sets forth standards for the formation and use of advisory committees.

SUMMARY: ACF and HRSA announce the renewal of the Advisory Committee on the Maternal, Infant and Early Childhood Home Visiting Program Evaluation to provide advice to the Secretary of Health and Human Services (''the Secretary'') on the design, plan, progress, and findings of the evaluation required under the Act.

FOR FURTHER INFORMATION CONTACT: T'Pring Westbrook, Administration for Children and Families; *tpring.westbrook@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: Sec. 511(g)(1) of Title V of the Social Security Act mandates an Advisory Committee to review, and make recommendations on, the design and plan for the evaluation required under the Act. To comply with the authorizing directive and guidelines under the Federal Advisory Committee Act (FACA), a charter has been filed with the Committee Management Secretariat in the General Services Administration (GSA), the appropriate committees in the Senate and U.S. House of Representatives, and the Library of Congress to establish the Advisory Board as a non-discretionary federal advisory committee. The charter was filed on January 27, 2015.

Objectives and Scope of Activitie

The purpose of the Committee is to provide advice and make recommendations to the Secretary of Health and Human Services, through the Assistant Secretary, ACF, and the Administrator, HRSA, with respect to the design, plan, progress, and results of the evaluation.

Membership and Designation

The Committee shall consist of up to 25 members appointed by the Secretary. Members shall be experts in the areas of program evaluation and research, education, and early childhood development. Members shall be appointed as Special Government Employees. The committee shall also include ex-officio members representing ACF, HRSA, and other agencies of the federal government designated by the Secretary as ex-officio members. The ACF Assistant Secretary and HRSA Administrator each shall recommend nominees for Co-Chairs of the Committee.

Members shall be invited to a 3-year term; such terms are contingent upon the renewal of the Committee by appropriate action prior to its termination.

Administrative Management and Support

Coordination, management, and operational services shall be provided by ACF, with assistance from HRSA.

A copy of the Committee charter can be obtained from the designated contact or by accessing the FACA database that is maintained by the GSA Committee Management Secretariat. The Web site for the FACA database is *http://fido.gov/facadatabase/.*

Dated: March 25, 2015.

Mary K. Wakefield,

Administrator, HRSA.

Mark H. Greenberg,

Acting Assistant Secretary, ACF.

[FR Doc. 2015–07978 Filed 4–6–15; 8:45 am]

BILLING CODE CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Tribal Consultation Meetings

AGENCY: Office of Head Start (OHS), Administration for Children and Families, HHS.

ACTION: Notice of meetings.

SUMMARY: Pursuant to the Improving Head Start for School Readiness Act of 2007, Public Law 110–134, notice is hereby given of three 1-day Tribal Consultation Sessions to be held between the Department of Health and Human Services, Administration for Children and Families, OHS leadership and the leadership of Tribal Governments operating Head Start (including Early Head Start) programs. The purpose of these Consultation Sessions is to discuss ways to better meet the needs of American Indian and Alaska Native children and their families, taking into consideration funding allocations, distribution formulas, and other issues affecting the delivery of Head Start services in their geographic locations [42 U.S.C. 9835, Section 640(l)(4)].

DATES:

June 16, 2015, from 1:00 p.m. to 5:00 p.m.;

July 30, 2015, from 1:00 p.m. to 5:00 p.m.;

August 17, 2015, from 1:00 p.m. to 5:00 p.m.

Locations:

- June 16, 2015—National Indian Head Start Directors Association, Hyatt