[BILLING CODE 4140-01-P]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-day comment request: Generic Clearance to Support the Safe to Sleep Campaign at the Eunice Kennedy Shriver National Institute for Child Health and Human Development (NICHD)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Child Health and Human Development, the National Institutes of Health, has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the Federal Register on December 30, 2013, pages 79472-79473 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institute of Child Health and Human Development, National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

DIRECT COMMENTS TO OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget,

Office of Regulatory Affairs, OIRA_submission@omb.eop.gov or by fax to 202-395-6974, Attention: NIH Desk Officer.

COMMENT DUE DATE: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

FOR FURTHER INFORMATION: To obtain a copy of the data collection plans and instruments, or request more information on the proposed project, contact: Dr. Sarah L. Glavin, Deputy Director, Office of Science Policy, Analysis and Communication, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 31 Center Drive, Room 2A18, Bethesda, Maryland 20892, or call a non-toll free number (301) 496-1877 or Email your request, including your address to glavins@mail.nih.gov. Formal requests for additional plans and instruments must be requested in writing.

PROPOSED COLLECTION: Generic Clearance to Support the Safe to Sleep Campaign at the Eunice Kennedy Shriver National Institute for Child Health and Human Development (NICHD), 0925-NEW, Eunice Kennedy Shriver National Institute of Child Health and Human Development (NICHD), National Institutes of Health (NIH).

Need and Use of Information Collection: This is a request for a new generic clearance that would be used for submissions specific to the Eunice Kennedy Shriver National Institute of Child Health and Human Development (NICHD) Safe to Sleep

(STS) public education campaign. Submissions for the STS campaign will be used to assess the understanding and reach of STS campaign materials and messages, and to monitor and improve campaign activities such as training workshops and overall implementation. The purpose of this information collection is to monitor and modify campaign activities, to plan future campaign activities, to develop messages and materials, and to develop distribution and outreach strategies that are effective at communicating their message to bring about the intended response, awareness, and/or behavioral change for the target audiences. This generic clearance will enable the NICHD to: 1) more efficiently assess the implementation of campaign activities; 2) better understand the target audiences' knowledge, attitudes, and beliefs toward STS messages and materials; 3) better understand how the campaign activities have influenced the target audiences' behaviors and practices; and 4) monitor and improve activities such as trainings, and material/message development. Having a way to gather feedback on the STS campaign activities is critical to assessing the reach and effect of campaign efforts. Data collected for the campaign can inform where future STS campaign resources can produce the most meaningful results.

Data collected for the STS campaign generic clearance will be used by a number of audiences, including STS campaign staff, NICHD leadership, STS campaign collaborators, Federal Sudden and Unexpected Infant Deaths (SUID)/Sudden Infant Death Syndrome (SIDS) Workgroup members, SUID/SIDS stakeholders, clinical and maternal/child health professionals, parents and caretakers, and the general public. These audiences may use the information collections to: 1) develop new campaign messages, materials, and/or training curricula; 2) monitor and improve campaign

activities; 3) make decisions about campaign activities; 4) inform current campaign activities; and 5) inform and/or change practices and behaviors of program participants.

Examples of the types of information collections that could be included under this generic clearance include: Focus groups and in-depth interviews with parents/caregivers and/or health professionals to get feedback on distribution and outreach activities, and/or campaign messages; and Surveys with parents/caregivers and/or health professionals to: 1) assess the usefulness of the new STS campaign materials, including print and on-line materials and a video, 2) track outreach experiences of program participants, 3) assess training participants' changes in knowledge related to safe infant sleep behavior and implementation of outreach methods taught, and 4) assess program participants' resource needs.

The sub-studies for this generic will be small scale, designed to obtain results frequently and quickly to guide campaign development and implementation, inform campaign direction, and be used internally for campaign management purposes. NICHD's current scope and capacity for STS generic sub-studies is non-existent and this request would fill this gap.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 3,000.

Estimated Annualized Burden Hours

Table 1: Estimates for Annual Burden Hours				
Type of Data Collection	Number of Respondents	Frequency of Response	Average Time per Response	Annual Hour Burden

Instrument				
Focus Groups	500	1	1	500
Pre/Post Test	2,500	1	15/60	625
Survey	2,500	1	15/60	625
Interview	500	1	1	500
Tracking/Feedback Form	1,500	1	30/60	750
Total	7,500			3,000

Dated: May 20, 2014

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Sarah L. Glavin, Ph.D.

Deputy Director, Office of Science Policy, Analysis, and Communications

Eunice Kennedy Shriver National Institute of Child Health and Human Development

National Institutes of Health

[FR Doc. 2014-12370 Filed 05/28/2014 at 8:45 am; Publication Date: 05/29/2014]