

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

**SUMMARY:** Under the provisions of the Paperwork Reduction Act of 1995, the Regulatory Secretariat Division 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 concerning Request for Authorization of Additional Classification and Rate, Standard Form (SF) 1444.

**DATES:** Submit comments on or before June 30, 2017.

**ADDRESSES:** Submit comments identified by Information Collection 9000-0089 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 9000-0089. Select the link “Comment Now” that corresponds with “Information Collection 9000-0089, Request for Authorization of Additional Classification and Rate, SF 1444.” Follow the instructions provided on the screen. Please include your name, company name (if any), and “Information Collection 9000-0089, Request for Authorization of Additional Classification and Rate, SF 1444” on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Sosa/IC 9000-0089.

*Instructions:* Please submit comments only and cite Information Collection 9000-0089, in all correspondence related to this collection. Comments received generally will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check [www.regulations.gov](http://www.regulations.gov), approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

**FOR FURTHER INFORMATION CONTACT:** Ms. Zenaida Delgado, Procurement Analyst, Federal Acquisition Policy Division, GSA, 202-969-7207 or email [zenaida.delgado@gsa.gov](mailto:zenaida.delgado@gsa.gov).

**SUPPLEMENTARY INFORMATION:**

**A. Purpose**

Federal Acquisition Regulation (FAR) 22.406 prescribes labor standards for federally financed and assisted

construction contracts subject to the Davis-Bacon and Related Acts (DBRA), as well as labor standards for non-construction contracts subject to the Contract Work Hours and Safety Standards Act (CWHSSA).

The recordkeeping requirements in this regulation, FAR 22.406, reflect the requirements cleared under OMB control numbers 1235-0023, 1235-0008, and 1235-0018 for 29 CFR 5.5(a)(1)(i), 5.5(c), and 5.15 (records to be kept by employers under the Fair Labor Standards Act (FLSA)). The regulation at 29 CFR 516 reflects the basic recordkeeping and reporting requirements for the laws administered by the Department of Labor Wage and Hour Division.

FAR 22.406-3, implements the recordkeeping and information collection requirements prescribed in 29 CFR 5.5(a)(1)(ii) cleared under OMB control number 1235-0023 (also prescribed at 48 CFR 22.406 under OMB control number 9000-0089), by providing SF 1444, Request for Authorization of Additional Classification and Rate, for the contractor and the Government to enter the recordkeeping and information collection data required by 29 CFR 5.5(a)(1)(ii) prior to transmitting the data to the Department of Labor.

This SF 1444 places no further burden on the contractor or the Government other than the information collection burdens already cleared by OMB for 29 CFR 5.

**B. Annual Reporting Burden**

*Number of Respondents:* 3,831.  
*Responses per Respondent:* 2.  
*Total Annual Responses:* 7,662.  
*Review time per response:* 5.  
*Total Burden Hours:* 3831.

**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:* Requester may obtain a copy of the justification from the General Services Administration, Regulatory Secretariat

Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202-501-4755. Please cite OMB Control No. 9000-0089, Request for Authorization of Additional Classification and Rate, SF 1444, in all correspondence.

Dated: April 25, 2017.

**Lorin S. Curit,**

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

[FR Doc. 2017-08670 Filed 4-28-17; 8:45 am]

**BILLING CODE 6820-EP-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-17-17AW]

**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](mailto: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**

Assessment of Targeted Training and Technical Assistance (TTA) Efforts on the Implementation of Comprehensive Cancer Control—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

The Centers for Disease Control and Prevention’s (CDC) National Comprehensive Cancer Control Program (NCCCP) has been a primary funder for state and community-based cancer control interventions since its inception in the late 1990s. In addition, CDC’s Office on Smoking and Health (OSH) also has worked to build state health department infrastructure and capacity to conduct coordinated comprehensive tobacco prevention and control activities which contribute to cancer health outcomes through the provision of funding to state health departments and local partners through the Nation State Based Tobacco Control Program (NSTB).

In striving to build capacity and maximize the impact of CDC’s funded programs, CDC has focused on developing and implementing innovative programs to enhance the training and technical assistance (TTA) delivered to NCCCP and NSBT grantee programs. CDC funds 10 organizations

under two cooperative agreements: The Consortium of National Networks to Impact Populations Experiencing Tobacco-Related and Cancer Health Disparities (DP13-1314), and National Support to Enhance Implementation of Comprehensive Cancer Control Activities (DP13-1315). Under these cooperative agreements, DP13-1314 and DP13-1315 awardees provide TTA to state NCCCP and NSBT grantees to support local implementation of high-impact public health strategies. Using two different TTA models, DP13-1314 and DP13-1315 aim to impact both short- and long-term outcomes on the awardee, NCCCP program, and population levels.

CDC proposes to conduct an assessment of the DP13-1314 and DP13-1315 cooperative agreements to: (1) Increase CDC’s understanding of the TTA provided to NCCCP and NSTB grantees across both cooperative agreements, (2) help identify the extent to which core elements of the TTA were administered, and (3) determine the elements of TTA across both cooperative agreements that show promise for improving NCCCP and NSTB capacity. There are no other data collection efforts currently underway to assess implementation of the two TTA models or their perceived effectiveness.

This information collection request will involve three complementary data collection efforts: (1) Case studies of DP13-1314 and DP13-1315 awardees (consisting of interviews with DP13-1314 and DP13-1315 program managers/directors, evaluators, and partners); (2) a cross-sectional web-based survey administered to NCCCP and NSBT program directors, coalition members, and partners; and (3) in-depth interviews with selected NCCCP and

NSBT program directors, staff, coalition members, and partners who received a high volume of TTA from one or more of the DP13-1314 and DP13-1315 awardees. The case studies will be used to explore how DP13-1314 and DP13-1315 awardees are implementing their respective cooperative agreements and administering TTA to NCCCP and NSBT grantees; the factors that affect the implementation of specific TTA components; and the extent to which each cooperative agreement was able to achieve planned short-term outcomes. The Web-based survey will inform CDC’s understanding of the reach of DP13-1314 and DP13-1315 TTA efforts; elicit information from NCCCP and/or NSBT programs and coalitions about the TTA received, including type, dosage, frequency and format; and assess the perceptions of the effectiveness of the TTA provided in building capacity to achieve intended outcomes. The in-depth interviews with “high-volume” TTA users will facilitate an in-depth exploration of the type and quality of TTA activities received; perceived quality of TTA and its contributions to NCCCP and NSBT grantee program implementation, and achievement of CDC priorities and goals.

CDC will use findings from the assessment to inform development of future TTA efforts that utilize the core elements across the two models to more effectively and efficiently support NCCCP’s partner organizations.

OMB approval is requested for 2 years. Participation is voluntary and respondents will not receive incentives for participation. There are no costs to respondents other than their time. The total estimated annualized burden hours are 231.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
DP13-1314 and DP13-1315 Awardee Organizations.	Worksheet for Identifying Case Study Interviewees.	5	1	1
DP13-1314 Program Directors/Managers .....	Case Study Interview Guide for DP13-1314 Program Directors/Managers.	4	1	1.5
	Case Study Follow-Up Interview Guide for DP13-1314 Program Directors/Managers.	4	1	1
DP13-1315 Directors/Managers .....	Case Study Interview Guide for DP1-1315 Program Directors/Managers.	1	1	1.5
	Case Study Follow-Up Interview Guide for DP1-1315 Program Directors/Managers.	1	1	1
DP13-1314 Evaluators .....	Case Study Interview Guide for DP1-1314 Evaluators.	4	1	1
DP13-1315 Evaluators .....	Case Study Interview Guide for DP1-1315 Evaluators.	1	1	1
DP13-1314 Partners .....	Case Study Interview Guide for DP1-1314 Partners.	8	1	1

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
DP13–1315 Partners .....	Case Study Interview Guide for DP1–1315 Partners.	2	1	1
NCCCP and NSBT Program Directors, Staff, Coalition Members, and Partners.	Web-based survey .....	780	1	15/60
NCCCP and NSBT Program Directors, Staff, Coalition Members, and Partners.	In-Depth Interview Guide .....	5	1	0.5

**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.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60Day–17–17ADR; Docket No. CDC–2017–0042]

**Proposed Data Collections Submitted for Public Comment and Recommendations**

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the Study to Explore Early Development, Teen Follow-Up Study (SEED Teen).

**DATES:** Written comments must be received on or before June 30, 2017.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC–2017–0042 by any of the following methods:

- *Federal eRulemaking Portal:* Regulations.gov. Follow the instructions for submitting comments.
- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

*Instructions:* All submissions received must include the agency name and Docket number. All relevant comments received will be posted without change to Regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to Regulations.gov.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: omb@cdc.gov.

**SUPPLEMENTARY INFORMATION:** Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 6501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of the information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of the information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget (OMB) approval. 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; (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; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information. Written comments should be received within 60 days of this notice.

**Proposed Project**

Study to Explore Early Development, Teen Follow-Up Study (SEED Teen)—New—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

Autism spectrum disorder (ASD) is a neurodevelopmental disorder characterized by impairments in social interaction and communication and stereotyped behaviors and interests. The U.S. prevalence of ASD is estimated at 1% to 2%. In addition to the profound, lifelong impacts on individuals’ functioning given the core deficits in social-communication abilities, a high proportion of children with ASD also have one or more other developmental impairments such as intellectual disability or attention-deficit-hyperactivity-disorder and children with ASDs have higher than expected