

multi-factorial fall prevention program to measure its impact on health outcomes for the elderly as well as acute and long-term care use and cost. The study is being conducted among a sample of individuals with private long-term care insurance who are age 75 and over using a multi-tiered random experimental research design to evaluate the effectiveness of the proposed fall prevention intervention program. The project began in Spring 2008 and is expected to be completed in December 2014.

*Need and Proposed Use of the Information:* The project will provide information to advance Departmental goals of reducing injury and improving the use of preventive services to positively impact Medicare use and spending.

*Likely Respondents:* Adults age 75 or older.

*Burden Statement:* Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. 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. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Forms (if necessary)	Type of respondent	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Initial Telephone Screen .....	Active Control Group (ACG)/Experimental Group (EG).	835	1	20 minutes ...	278
In-person interview .....	EG .....	435	1	1.25 hours ...	544
Jump start phone call .....	EG .....	435	1	30 minutes ...	218
Quarterly phone calls .....	ACG/EG .....	835	4	10 minutes ...	556
Final Telephone Screen .....	ACG/EG .....	167	1	20 minutes ...	56
Final In-person interview .....	EG .....	167	1	1.25 hours ...	209
Total .....	.....	.....	.....	.....	1861

**Keith A. Tucker,**  
*Information Collection Clearance Officer.*  
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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-13-13EP]

**Agency Forms Undergoing Paperwork Reduction Act Review**

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639-7570 or send an email to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

**Proposed Project**

Million Hearts® Hypertension Control Challenge—New—National Center for

Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

CDC proposes to launch the Million Hearts® Hypertension Control Challenge to identify clinical practices and health systems that have been successful in achieving high rates of hypertension control and to develop models for dissemination. The most successful clinical practices or health plans will be recognized as Million Hearts® Hypertension Control Champions.

CDC requests OMB approval to collect the information needed to identify, qualify, and rank applicants for recognition through the Million Hearts® Hypertension Control Challenge. Interested providers or clinical programs voluntarily self-nominate their practice or healthcare system by completing a web-based nomination form located on the Challenge.gov web portal. The nomination process will include submission of the minimum amount of data needed to provide evidence of clinical success in achieving hypertension control, including: (a) Two point-in-time measures of the clinical hypertension control rate for the patient population, (b) the size of the clinic population served, and (c) a description

of the sustainable systems adopted to achieve hypertension control rates.

CDC scientists or contractors will assign a preliminary score to each submitted nomination form. Those with the highest preliminary scores will be further reviewed by a CDC-sponsored panel of three to five experts in hypertension control. The panel will provide CDC with a ranked list of nominees.

Finalists will be asked to participate in a data verification process that includes verification of how information was obtained from electronic records, remote electronic record or chart review, on-site review, or verification with other sources. Finalists may be eliminated based on the results of data verification.

Each remaining finalist, or Champion, will be asked to participate in a semi-structured interview. The interview will provide detailed information about the strategies employed by the practice or health system to achieve exemplary rates of hypertension control, including barriers and facilitators for those strategies.

OMB approval is requested for three years to support three annual Challenges.

There are no costs to respondents other than their time. The total estimated burden hours are 958.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs)
Physicians (Single or Group Practices or Health System).	Million Hearts® Hypertension Control Champion Nomination Form.	1,735	1	.5
Finalists .....	Million Hearts® Hypertension Control Champion Data Verification Form.	30	1	1
Selected Champion .....	Interview Guide: Million Hearts® Hypertension Control Champion.	30	1	2

**Ron A. Otten,**

Director, 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**

[30Day-13-0915]

**Agency Forms Undergoing Paperwork Reduction Act Review**

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639-5960 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov). Send written comments to 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 to Support the Development of Sickle Cell Disease

Educational Messages and Materials for the Division of Blood Disorders (0920-0915, Expired 01/31/2013)—Reinstatement—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

CDC seeks to improve the quality of life of people living with sickle cell disease (SCD). To accomplish this goal, CDC aims to address the need for educational messages and materials for adolescents, young adults, adults, and older adults living with SCD. CDC is interested in understanding the informational needs of these audiences related to the adoption of healthy behaviors and the prevention of complications associated with sickle cell disease. To develop valuable messages and materials, CDC will conduct formative focus groups with people with SCD across the country. Participants will stem from four urban centers as well as more remote, rural areas. Based on the findings from the formative focus groups, CDC will develop and test draft messages.

A total of 10 focus groups will be conducted. Eight focus groups with people with SCD would be held in four cities: Atlanta, GA; Detroit, MI; Oakland, CA; and Philadelphia, PA. Two in-person focus groups—one with males and one with females—will be

conducted in each city with each target audience: adolescents aged 15-17, young adults aged 18-25, adults aged 26-35, and older adults 36 and over. To reach more rural participants, two telephone focus groups will be conducted: one with female adolescents aged 15-17 and a second with male older adults aged 36 and older.

The focus groups will be conducted with eight to nine participants in each and will last 2 hours. As part of the focus group, participants will complete an informed consent or adolescent assent form before discussion begins. The parents of the expected 27 adolescent participants (three groups of 9 each) will fill out a permission form to provide their consent in advance of the groups. The use of trained moderators and a structured moderator's guide will ensure that consistent data are collected across the groups. In total, up to 90 people with SCD will participate in the focus group data collection. It is estimated that 120 potential participants will need to be screened to reach the target of 90 participants. The estimated time per response for screening and recruitment is 12 minutes.

CDC requests OMB approval to extend clearance for one year. There is no cost to respondents other than their time. The estimated annualized burden hours for this data collection activity are 204.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hours)
Parents of adolescents (aged 15-17) living with SCD. Young adults (aged 18-25) living with SCD. Adults (aged 26-35) living with SCD. Older adults (aged 36+) living with SCD.	Participant Screener and Recruitment Script	120	1	12/60
Adolescents (aged 15-17) living with SCD .... Young adults (aged 18-25) living with SCD. Adults (aged 26-35) living with SCD. Older adults (aged 36+) living with SCD.	Focus Group Moderator's Guide .....	90	1	2