

survey to assess their involvement in promoting HIV education, awareness, and policies in their organization. (e) Partnerships Activities Form: Partners may be asked to complete a brief electronic form to provide information on each partner activity that they complete. The form will collect information on information such as the type of event, the audience, and key

highlights; the number of HIV tests administered (if any) and the number of preliminary positives; the number and type of materials distributed. This information will allow CDC to know what partners are doing to advance HIV prevention and education, and how CDC can alter their partnership efforts to facilitate HIV prevention and education in the future.

The information obtained from the proposed study will be used by federal policy makers to assess the effectiveness of the partnership activities and the appropriateness of continued or expanded funding of partnership projects.

There is no cost to participants other than their time. The total estimated annualized burden hours are 5,200.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Partner Organization	Quarterly Metric Database	50	4	18	3,600
Partner Organization	Bi-annual Key Informant Interview ..	25	2	1	50
Partner Organization	Interim Progress Reports	25	2	8	400
Partner Organization	Partner Survey	300	1	30/60	150
Partner Organization	Partnerships Activities Form	500	4	25/60	1,000
Total	5,200

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

[30Day-16-16CO]

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

Developing a Self-Management Tool for Individuals with Systemic Lupus Erythematosus—New—National Center for Chronic Disease Preventions and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Systemic Lupus Erythematosus (SLE) is an autoimmune disease in which the immune system produces antibodies to cells within the body leading to widespread inflammation and tissue damage. SLE has a variety of clinical

manifestations and can affect joints, skin, the brain, lungs, kidneys, and blood vessels.

Effective SLE management depends not only upon clinical interventions, but also on self-management— those things done on a day-to-day basis to manage SLE. SLE self-management requires gaining essential knowledge, skills, and confidence to manage the condition.

CDC previously launched a two-year project called “Filling a Gap: Creating Educational Program, Tools, or Materials to Enhance Self-Management in Systemic Lupus Erythematosus” to identify and address the needs of lupus patients in practicing effective self-management. The purpose of this project is to develop a SLE self-management tool to improve the ability of people living with lupus to manage their condition.

The proposed information collection will assess a prototype CDC SLE self-management tool that is in development to ensure that the tool is usable and useful to members of the target audience. The tool is expected to be comprised of multiple SLE self-management resources that may include, but are not limited to: Education resources about fatigue management, pain management, healthy diet, and exercise; symptom trackers; medication trackers; appointment calendars; resources about communication with family, friends, and co-workers about SLE; and strategies for coping with depression and anxiety.

CDC plans to make the tool available in an electronic format (web-based or a native mobile application) and will

consider making it available as a printed resource, depending on the feedback obtained during the testing process.

The information collection will also gauge the needs of the target audience(s), tool format and delivery method(s), and the tool's clarity, relevance, salience and appeal. A series of focus groups with women with a diagnosis of SLE, and one-on-one telephone interviews with men with a

diagnosis of SLE will be conducted to assess the tool. The same discussion guide will be used for all information collections.

The estimated burden per response for participating in a focus group discussion is two hours. The estimated burden per response for a discussion conducted via telephone interview is 45 minutes. Respondent burden also includes two hours for reviewing the

prototype CDC SLE Self-management Tool in advance of the focus group meeting or telephone interview.

OMB approval is requested for one year. Participation is voluntary and there are no costs to respondents other than their time.

The total estimated burden hours are 646.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Women with SLE diagnosis	Screener for Women	192	1	10/60
	Prototype CDC SLE Self-management Tool	128	1	2
	Discussion Guide for Use in Focus Groups with Women or Interviews with Men.	128	1	2
Men with SLE diagnosis	Screener for Men	40	1	10/60
	Prototype CDC SLE Self-management Tool	20	2	2
	Discussion Guide for Use in Focus Groups with Women or Interviews with Men.	20	1	45/60

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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day-16-0017; Docket No. CDC-2016-0014]

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 efforts 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 proposed revisions of the information collection entitled *Application for Training* (OMB Control No. 0920-0017). CDC seeks to request

Office of Management and Budget approval to (1) continue to collect information through the use of the Training and Continuing Education Online New Participant Registration form for new learners to establish an account that provides CDC necessary information to process learner requests for continuing education, and (2) implement a new electronic information collection through the use of the Training and Continuing Education Online Proposal form that allows training developers to provide CDC necessary information to process and accredit trainings for continuing education.

DATES: Written comments must be received on or before April 1, 2016.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2016-0014 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.

Please note: All public comment should be submitted through the Federal eRulemaking portal (Regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION 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.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of 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 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 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