

ESTIMATED ANNUALIZED BURDEN TABLE—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden (in hrs.)
ADSC Director/Designated Staff	ADSC Questionnaire	2,720	1	30/60	1,360
RCC and ADSC Directors/Designated Staff.	Data Retrieval	429	1	5/60	36
Total	4,319

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. 2015-28475 Filed 11-9-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-16-16CO; Docket No. CDC-2015-
0099]

Proposed Data Collection 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 for Developing a Self-
Management Tool for Individuals with
Systemic Lupus Erythematosus (SLE), to
assess the value of a tool aimed to
enhance the ability of persons with SLE
to effectively manage their condition.

DATES: Written comments must be
received on or before January 11, 2016.

ADDRESSES: You may submit comments,
identified by Docket No. CDC-2015-
0099 by any of the following methods:

Federal eRulemaking Portal:
Regulation.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: To
request more information on the
proposed project or to obtain a copy of
the information collection plan and
instruments, contact the 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. 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
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.

Proposed Project

Developing a Self-Management Tool
for Individuals with Systemic Lupus
Erythematosus (SLE)—New—National
Center for Chronic Disease Prevention
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 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 collection. The estimated burden per response for participating in a focus group discussion is 2 hours. The estimated burden per response for a discussion conducted via telephone interview is 45 minutes. Respondent burden also includes 2 hours for reviewing the draft 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.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden (in hrs.)
Women with SLE diagnosis.	Screener	192	1	10/60	32
	Review of the CDC SLE Self-Management Tool	128	1	2	256
	Discussion Guide	128	1	2	256
Men with SLE diagnosis	Screener	40	1	10/60	7
	Review of the CDC SLE Self-Management Tool	20	2	2	40
	Discussion Guide	20	1	45/60	15
Total	606

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–16–16CP; Docket No. CDC–2015–0100]

Proposed Data Collection 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 a proposed information collection request entitled “Data Collection for Community-based Tick Control for the Prevention of Rocky Mountain Spotted Fever in Hermosillo, Mexico.” This project will be carried out in collaboration with the Rickettsial Zoonoses Branch, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC) and the University of Sonora School of Medicine (UNSOM) to assess the efficacy and impact of a community based tick prevention project.

DATES: Written comments must be received on or before January 11, 2016.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2015–0100 by any of the following methods:

- Federal eRulemaking Portal: [Regulations.gov](http://www.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](http://www.Regulations.gov), including any personal information provided. For access to the docket to read background documents or comments received, go to [Regulations.gov](http://www.Regulations.gov).

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

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 the 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. 3501–3520), Federal agencies must obtain approval from the Office of