

Dated: March 31, 2009.

Bruce G. Gellin,

*Deputy Assistant Secretary for Health,
Director, National Vaccine Program Office,
U.S. Department of Health and Human
Services.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-09-08AU]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-5960 and send comments to Maryam Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta,

GA 30333 or send an e-mail to omb@cdc.gov.

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; and (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. Written comments should be received within 60 days of this notice.

Proposed Project

Assessing Problem Areas in Referrals for Chronic Hematologic Malignancies and Developing Interventions to Address Them—New—Division of Cancer Prevention and Control, National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

One of the six aims of the Institute of Medicine's Crossing the Quality Chasm report is to improve the timeliness of care for patients. Data from Europe and

Canada, as well as single-site studies in the United States, allude to a problem of timely referral and diagnosis for patients with cancer. Despite the advent of new diagnostics and therapeutics for patients with chronic hematological malignancies, the size and scope of a potential problem regarding their referral from primary care providers to specialists is not well-defined in the current literature.

CDC proposes to conduct a one-time study to collect qualitative and quantitative information on optimal and sub-optimal referral patterns for patients with confirmed or suspected chronic hematologic malignancies. Information will be collected to identify specific factors related to delays in diagnosis and/or referral to appropriate medical specialists. Information will be collected through in-depth interviews with hematologic cancer patients, in-depth interviews and focus groups with primary care providers, interviews with specialists in hematology and oncology in Texas, and a one-time postal survey to a sample of primary care providers in Massachusetts.

The ultimate goal is to develop tools that will improve the awareness, diagnosis, and referral of persons with chronic hematological cancers by primary care providers.

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	Avg. burden per response (in hrs)	Total burden (in hours)
Community Hematologists and Oncologists.	Hematologists and Oncologists Interview Phone Recruitment Script.	100	1	2/60	3
	Hematologists and Oncologists Interview Guide.	18	1	1.5	27
Patients	Patient Interview Phone Recruitment Script.	50	1	2/60	2
Primary Care Providers (PCP)	Patient Interview Guide	18	1	1.5	27
	PCP Survey Cover Letter	250	1	2/60	8
	PCP Survey	150	1	20/60	50
	PCP Opt-Out Card	100	1	2/60	3
	PCP Survey Reminder Letter	200	1	2/60	7
	PCP Interview Phone Recruitment Script.	100	1	3/60	5
	PCP Interview Guide	18	1	1.5	27
	PCP Focus Group Phone Recruitment Script.	50	1	3/60	3
Total	PCP Focus Group Guide	18	1	2	36
					198

Dated: April 6, 2009.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E9-8337 Filed 4-10-09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Aging

Agency Information Collection Activities; Proposed Collection; Comment Request; SMP (Formerly Senior Medicare Patrol) Program Outcome Measurement

AGENCY: Administration on Aging, HHS.

ACTION: Notice.

SUMMARY: The Administration on Aging (AoA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to SMP (formerly Senior Medicare Patrol) Program outcome measurement.

DATES: Submit written or electronic comments on the collection of information by June 12, 2009.

ADDRESSES: Submit electronic comments on the collection of information to:

doris.summey@aoa.hhs.gov.

Submit written comments on the collection of information to Administration on Aging, Washington, DC 20201. Attention: Doris Summey.

FOR FURTHER INFORMATION CONTACT:

Doris Summey, by telephone: (202) 357-3533 or by e-mail:

doris.summey@aoa.hhs.gov.

SUPPLEMENTARY INFORMATION: Under the 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. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in

the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

Grantees are required by Congress to provide information for use in program monitoring and for Government Performance and Results Act (GPRA) purposes. This information collection reports the number of active volunteers, issues and inquiries received, other SMP program outreach activities, and the number of Medicare dollars recovered, among other SMP Performance outcomes.

AoA estimates the burden of this collection of information as follows: *Respondents:* 54 SMP grantees at 23 hours per month (276 hours per year, per grantee). *Total Estimated Burden Hours:* 14,904 hours per year.

Dated: April 8, 2009.

Edwin L. Walker,

Acting Assistant Secretary for Aging.

[FR Doc. E9-8386 Filed 4-10-09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-09-09BK]

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Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-5960 and send comments to Maryam I. Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an e-mail to *omb@cdc.gov*.

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; and (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. Written comments should be received within 60 days of this notice.

Proposed Project

Registration of Individuals Displaced by the Hurricanes Katrina and Rita (Pilot Project)—New—Agency for Toxic Substances and Disease Registry (ATSDR), Coordinating Center for Environmental Health and Injury Prevention (CCEHIP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

On August 29, 2005, Hurricane Katrina made landfall on the coast of the Gulf of Mexico near New Orleans, Louisiana, and became one of the most deadly and destructive storms in U.S. history. Also occurring in 2005, Hurricane Rita was the fourth-most intense Atlantic hurricane ever recorded and the most intense tropical cyclone ever observed in the Gulf of Mexico. Following the initial phase of the response, the Federal Emergency Management Agency (FEMA) assumed the primary role for housing displaced persons over the intermediate term. To support those needing temporary housing, FEMA provided over 143,000 travel trailers, park homes, and mobile homes for persons displaced by the above mentioned storms. However, some persons living in trailers complained of an odor or of eye or respiratory tract irritation.

FEMA entered into an Interagency Agreement with the Centers for Disease Control and Prevention (CDC)/ATSDR on August 16, 2007 to conduct a