

John D. Graubert,
Acting General Counsel.
[FR Doc. 06–5631 Filed 6–22–06; 8:45 am]
BILLING CODE 6750–01–C

**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

[60Day-06–0603]

**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 Seleda Perryman, CDC Assistant 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

Information Network (REACH IN)—Extension (0920–0603)—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Racial and Ethnic Approaches to Community Health 2010 (REACH 2010) currently funds forty local coalitions to establish community based programs and culturally appropriate interventions to eliminate racial and ethnic health disparities. Two previously funded grantees also retain access to the system. Communities served by REACH 2010 include: African American, American Indian, Hispanic American, Asian American, and Pacific Islander. These communities can select among infant mortality, deficits in breast and cervical cancer screening and management, cardiovascular diseases, diabetes, HIV/ AIDS, and deficits in childhood and adult immunizations to be the focus of their interventions. Guided by logic

models, each community articulates goals, objectives, and related activities; tracks whether goals and objectives are met, ongoing, or revised; and evaluates all program activities. This information is then entered into the REACH Information Network (REACH IN). REACH IN is a customized Internet-based support system that allows REACH 2010 grantees to perform remote data entry and retrieval of data.

This support system is designed to create on-demand graphs and reports of grantees’ activities and accomplishments, monitor progress toward the achievement of goals and objectives, and share and synthesize information across grantees’ activities. Both quantitative and qualitative analyses can be performed. These analyses relate primarily to three stages of the REACH 2010 logic model: Capacity building, targeted actions (interventions), and community and systems change and change among change agents. Users are supported with technical assistance and training, covering the usage of the system from a content/project goals perspective, and technical operations.

The annualized estimated burden is based on 42 respondents, including 40 currently funded grantees and two that were funded previously who retain access to the system. It is estimated that they each use the system four times a year to enter data, each data entry taking about 30 minutes. There are no costs to respondents except their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden hours
REACH 2010 grantees	42	4	30/60	84

Dated: June 15, 2006.
Joan F. Karr,
Acting Reports Clearance Officer, Centers for Disease Control and Prevention.
[FR Doc. E6–9919 Filed 6–22–06; 8:45 am]
BILLING CODE 4163–18–P

**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

[60Day–06–0214]

**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 Seleda Perryman, CDC Assistant 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

National Health Interview Survey (NHIS) 2007–2009, (OMB No. 0920–0214)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k) authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on the extent and nature of illness and disability of the population of the United States.

The annual National Health Interview Survey is a major source of general

statistics on the health of the U.S. population and has been in the field every year since 1957. This household-based survey collects demographic and health-related information on a nationally representative sample of households throughout the country. The survey has three modules: The family module collects information on everyone in the family; the sample adult module collects more detailed information on a randomly selected adult; and the sample child module collects information on a randomly selected child (in households with children). Information is collected using computer assisted personal interviews (CAPI). A core set of data is collected each year while sponsored supplements vary from year to year. In addition to the core data collection, in 2007 there will be two new supplements, which will provide additional data on complementary and alternative medicine (including questions on topics such as acupuncture, chiropractic or osteopathic manipulation, meditation, natural herbs, and yoga) and on hearing disorders (such as hearing loss and

tinnitus). These supplements are sponsored by the National Center on Complementary and Alternative Medicine and the National Institute on Deafness and Other Communication Disorders, both parts of the National Institutes of Health.

In accordance with the 1995 initiative to increase the integration of surveys within the Department of Health and Human Services, respondents to the NHIS serve as the sampling frame for the Medical Expenditure Panel Survey conducted by the Agency for Healthcare Research and Quality. The NHIS has long been used by government, university, and private researchers to evaluate both general health and specific issues, such as cancer, diabetes, and access to health care. It is a leading source of data for the Congressionally-mandated “Health US” and related publications, as well as the single most important source of statistics to track progress toward the National Health Promotion and Disease Prevention Objectives, “Healthy People 2010.”

There is no cost to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden hours
Family member	39,000	1	21/60	13,650
Sample adult	32,000	1	42/60	22,400
Sample child	13,000	1	15/60	3,250
Total				39,300

Dated: June 16, 2006.

Joan F. Karr,

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

[FR Doc. E6–9920 Filed 6–22–06; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–296]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid

Services (CMS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the Agency’s function; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Home Health Advance Beneficiary Notice (HHABN)

and Supporting Regulations in 42 CFR 411.404 and 484.10(a) and (e).; *Use:* Home health agencies (HHAs) are required to provide written notice to Medicare beneficiaries under various circumstances involving the initiation, reduction, or termination of services. The notice is designed to ensure that beneficiaries receive complete and useful information to enable them to make informed consumer decisions. Consistent with the decision of the U.S. Court of Appeals (2nd Circuit) in the *Lutwin v. Thompson*, HHAs must now also issue HHABNs in a broader set of circumstances in conjunction with their responsibilities under the Home Health Conditions of Participation (HH COPs). The HHABN instructions explain when the newly revised HHABN should be issued, and include additional changes to simplify notice policy for HHAs. The notice must be issued timely and provide clear and accurate information about the specified services and, if