hypertension, Type 2 diabetes, stroke, coronary heart disease, and osteoarthritis. Beyond the human costs, economic costs are extreme and are climbing. One estimate is that medical expenses related to this epidemic accounted for 9.1% of total U.S. medical expenditures in 1998, and the U.S. Surgeon General has estimated that direct and indirect costs related to obesity totaled $117 billion in 2000. Healthy People 2010 established goals for obesity reduction, which included targets of weight reduction of 15% for adults and 5% for children and youth.

Targeting communities at risk of overweight and obesity is an essential step toward realizing the goal of reversing current trends in obesity. Community-based programs to reduce risk of heart disease provide some models; however, outcomes vary and are affected by several confounding conditions. A report on prevention of childhood obesity, prepared by the Institute of Medicine in 2007, concluded that there are insufficient studies to generate recommendations for best practices in obesity prevention. Instead, the report compiles promising practices, including those set in communities.

CDC plans to apply methodology recommended by the CDC Task Force on Community Preventive Services to improve the translation and dissemination of promising practices into community-based obesity prevention programs. Information necessary to this purpose will be collected from the general public by a contractor. Information will be collected concerning respondents’ knowledge, attitudes, and beliefs about obesity and physical activity; the need for community leaders to encourage healthier diets and more physical activity; and opportunities for leveraging current community efforts.

Two hundred fifty respondents will be recruited to participate in four on-line, small-group discussions over a period of about one month. The discussions will utilize Voice over Internet Protocol technology and will be facilitated by a moderator. Each discussion will last one hour. In preparation for the initial discussion, respondents will receive a confirmation e-mail and will be asked to review a guide to on-line discussion groups. In addition, discussion group participants will be asked to review a set of briefing materials prior to the first on-line group meeting.

### ESTIMATED ANNUALIZED BURDEN HOURS

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<th>Type of respondents</th>
<th>Form name</th>
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<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
<th>Total burden (in hours)</th>
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Date: May 4, 2010.

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

[FR Doc. 2010–11060 Filed 5–7–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day–10–0743]

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 or send comments to Maryam I. Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS D–74, 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
to increasing overall rates, a significant public health priority in the U.S. is to reduce variation in breastfeeding rates across population subgroups. For example, in 2005, nearly three-quarters of white mothers started breastfeeding, but only about half of black mothers did so.

The health care system is one of the most important and effective settings to improve breastfeeding. In 2007, CDC conducted the first national survey of Maternity Practices in Infant Nutrition and Care (known as the mPINC Survey) in health care facilities (hospitals and free-standing childbirth centers). This survey was designed to provide baseline information and to be repeated every two years. The survey was conducted again in 2009. The survey inquired about patient education and support for breastfeeding throughout the maternity stay as well as staff training and maternity care policies.

Prior to the fielding of the 2009 iteration, CDC was requested to provide a report to OMB on the results of the 2007 collection. In this report, CDC provided survey results by geographic and demographic characteristics and a summary of activities that resulted from the survey.

Because the 2011 mPINC survey repeats the prior iterations (2007 and 2009), the methodology, content, and administration of it will match those used before. The census design does not employ sampling methods. Facilities are identified by using the American Association of Birth Centers (AABC) and the American Hospital Association (AHA) Annual Survey of Hospitals. In addition to all facilities that participated in 2007 or 2009, the 2011 survey will include those that were invited but did not participate in 2007 or 2009 and any that are new since then. All birth centers and hospitals with ≥1 registered maternity bed will be screened via a brief phone call to assess their eligibility, identify additional locations, and identify the appropriate point of contact. The extremely high response rates to the 2007 mPINC survey of 82 percent and 81 percent to the 2009 iteration indicate that the methodology is appropriate and also reflects unusually high interest among the study population.

As with the initial surveys, a major goal of the 2011 follow-up survey is to be fully responsive to their needs for information and technical assistance. CDC will provide direct feedback to respondents in a customized benchmark report of their results and identify and document progress since 2007 on their quality improvement efforts. National and state reports will use de-identified data to describe incremental changes in practices and care processes over time at the facility, state, and national levels.

Participation in the survey is voluntary, and responses may be submitted by mail or through a Web-based system. There are no costs to respondents other than their time.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; The Framingham Heart Study (FHS)

SUMMARY: 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 National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: The Framingham Heart Study. Type of Information Request: Revision (OMB No. 0925–0216). Need and Use of Information Collection: The Framingham Heart Study will conduct examinations and morbidity and mortality follow-up for the purpose of studying the determinants of cardiovascular disease. Examinations will be conducted on the original,