

service programs are both effective and efficient in influencing workplace change. This requires a social marketing evaluation of NIOSH products to measure the degree of customer satisfaction and their adoption of recommended actions.

The present project proposes to do this by conducting a survey of a primary segment of NIOSH's customer base, the community of occupational safety and health professionals. In collaboration with the American Association of Occupational Health Nurses (13,000 members), the American Industrial Hygiene Association (12,400 members),

the American College of Occupational and Environmental Medicine (6,500 members), and the American Society of Safety Engineers (33,000 members), NIOSH will survey a sample of their memberships to ascertain, among other things: (1) Their perceptions and attitudes toward NIOSH as a general information resource; (2) their perceptions and attitudes about specific types of NIOSH publications (e.g., criteria documents, technical reports, alerts); (3) the frequency and nature of referral to NIOSH in affecting occupational safety and health practices and policies; (4) the extent to which

they have implemented NIOSH recommendations; and (5) their recommendations for improving NIOSH products and delivery systems. The results of this survey will provide an empirical assessment of the impact of NIOSH publications on occupational safety and health practice and policy in the United States as well as provide direction for shaping future NIOSH communication efforts. Respondents will have the option of responding by mail or electronically through the NIOSH Web site. There is no costs to respondents for participation.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs.)	Total burden in hours
Survey Sample	600	1	20/60	200
American Association of Occupational Health Nurses				
American Industrial Hygiene Association				
American College of Occupational and Environmental Medicine				
American Society of Safety Engineers				
Total				200

Dated: December 20, 2002.

Nancy E. Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-03-30]

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 the CDC Reports Clearance Officer on (404) 498-1210.

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. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project: 2004 National Health Interview Survey.(0920-0214)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

The annual National Health Interview Survey (NHIS) is a basic source of general statistics on the health of the U.S. population. 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. This survey is 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, AIDS, and childhood immunizations. Journalists use its data

to inform the general public. It will continue to be 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."

Because of survey integration and changes in the health and health care of the U.S. population, demands on the NHIS have changed and increased, leading to a major redesign of the annual core questionnaire, or Basic Module, and a shift from paper questionnaires to computer assisted personal interviews (CAPI). These redesigned elements were partially implemented in 1996 and fully implemented in 1997. This clearance is for the eighth full year of data collection using the core questionnaire on CAPI, for the implementation of a supplement on children's mental health, and for a software field test to evaluate a switch from CASES software to Blaise software for the CAPI instrument. The field test for the new software is scheduled for June 2003. The data collection for the full survey is planned for January-December 2004, and will result in publication of new national estimates of health statistics, release of public use micro data files, and a sampling frame for other integrated surveys. There is no cost to the respondents other than their time.

SOFTWARE FIELD TEST IN JUNE 2003

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hours)	Total burden (in hours)
Family	300	1	21/60	105
Sample Adult	246	1	42/60	172
Sample Child	100	1	15/60	25
Total				302

FULL SURVEY JANUARY–DECEMBER 2004

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hours)	Total burden (in hours)
Family	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

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Nancy E. Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–03–26]

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Proposed Project: National Program of Cancer Registries—Cancer Surveillance System 0920–0469—Extension—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

The American Cancer Society estimates that about 1.2 million Americans will be newly diagnosed with cancer and that about 8.2 million Americans are currently alive with a history of cancer. The National Institutes of Health estimates the cost of cancer is about \$172 billion including (\$61 billion) direct costs to treat cancer and (\$111 billion) indirect costs in lost productivity due to illness and premature death.

In 2000, CDC implemented the National Program of Cancer Registries (NPCR)—Cancer Surveillance System (CSS) to collect, evaluate and disseminate cancer incidence data

collected by population-based cancer registries. In 2002, CDC published United States Cancer Statistics—1999 Incidence which provided cancer statistics for 78% of the United States population from all cancer registries whose data met national data standards. Prior to this, at the national level, cancer incidence data were available for only 14% of the population of the United States.

With this expanded coverage of the U.S. population, it will now be possible to better describe geographic variation in cancer incidence throughout the country and provide incidence data on minority populations and rare cancers to further plan and evaluate state and national cancer control and prevention efforts.

Therefore, the CDC's NCCDPHP, Division of Cancer Prevention and Control, proposes to continue to aggregate existing cancer incidence data from states funded by the National Program of Cancer Registries into a national surveillance system.

These data are already collected and aggregated at the state level. Thus the additional burden on the states is small. Funded states are asked to continue to report data to CDC on an annual basis twelve months after the close of a diagnosis year and again at twenty-four months to obtain more complete incidence data and vital status from mortality data. The estimated annualized cost to respondents is \$885,000.