

allowed the monitoring of progress toward these goals. For example, recipient efforts to establish public health–health care collaboration has been integrated into CDC’s 6|18 initiative which connects healthcare purchasers, payers, and providers with CDC researchers, economists, and policy analysts to find ways to improve health and control costs with the 6|18 interventions;

- Allow the NACP and the state asthma programs to make more informed decisions about activities to achieve objectives. For example, PM information identified a problem with enrolling patients most in need of intervention into recipient programs. This led to cross-state discussions and changes in recruitment strategies;
- Motivate use of data and evaluation findings. For example, the requirement

to report actions taken based on evaluation findings encourages program managers and health departments officials to act on recommendations in evaluation reports.

CDC requests OMB approval for an estimated 105 annual burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Funded Asthma Program Recipients	Performance Measures Reporting Tool.	30	1	150/60	75
	Emergency Department Visits Reporting Form.	30	1	30/60	15
	Hospital Discharge Reporting Form	30	1	30/60	15
Total .....	.....	.....	.....	.....	105

**Jeffrey M. Zirger,**

*Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.*

[FR Doc. 2022–27506 Filed 12–19–22; 8:45 am]

**BILLING CODE 4163–18–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day–23–1030]

**Agency Forms Undergoing Paperwork Reduction Act Review**

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Developmental Studies to Improve the National Health Care Surveys” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on September 30, 2022 to obtain comments from the public and affected agencies. CDC did not receive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th

Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

**Proposed Project**

Developmental Studies to Improve the National Health Care Surveys (OMB Control No. 0920–1030, Exp. 06/30/2023)—Extension—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), as amended, authorizes the Secretary of Health and Human Services (DHHS), acting through the Division of Health Care Statistics (DHCS) within NCHS, shall collect statistics on the extent and nature of illness and disability of the population of the United States.

The DHCS conducts the National Health Care Surveys (NHCS), a family of nationally representative surveys of encounters and health care providers in inpatient, ambulatory, and long-term care settings. This information collection request (ICR) is for the extension of a generic clearance to conduct developmental studies to improve this family of surveys. This three-year clearance period will include studies to evaluate and improve upon existing survey design and operations, as well as to examine the feasibility of, and address challenges that may arise with, future expansions of the National Health Care Surveys.

Specifically, this request covers developmental research with the

following aims: (1) to explore ways to refine and improve upon existing survey designs and procedures; and (2) to explore and evaluate proposed survey designs and alternative approaches to data collection. The goal of these research studies is to further enhance DHCS existing and future data collection protocols to increase research capacity and improve health care data quality for the purpose of monitoring public health and well-being at the national, state and local levels, thereby informing the health policy decision-making process. The information collected through this Generic ICR will not be used to make generalizable statements about the population of interest or to inform public policy; however, methodological findings may be reported.

This Generic ICR would include studies conducted in person, via the telephone or internet, and by postal or electronic mail. Methods covered would include qualitative (e.g., usability testing, focus groups, ethnographic studies, and respondent debriefing questionnaires) and/or quantitative (e.g., pilot tests, pre-tests and split sample experiments) research methodologies. Examples of studies to improve existing survey designs and procedures may include evaluation of incentive approaches to improve recruitment and increase participation rates; testing of new survey items to obtain additional data on providers, patients, and their encounters while minimizing misinterpretation and human error in data collection; testing data collection in panel surveys; triangulating and validating survey responses from multiple data sources; assessment of the feasibility of data retrieval; and development of protocols that will locate, identify, and collect accurate survey data in the least labor-intensive and burdensome manner at the sampled practice site.

To explore and evaluate proposed survey designs and alternative approaches to collecting data, especially with the nationwide adoption of

electronic health records, studies may expand the evaluation of data extraction of electronic health records and submission via continuity of care documentation to small/mid-size/large medical providers and hospital networks, managed care health plans, prison-hospitals, and other inpatient, outpatient, and long-term care settings that are currently either in-scope or out-of-scope of the National Health Care Surveys. Research on feasibility, data quality and respondent burden also may be carried out in the context of developing new surveys of health care providers and establishments that are currently out-of-scope of the National Health Care Surveys.

Specific motivations for conducting developmental studies include: (1) within the National Ambulatory Medical Care Survey (NAMCS), new clinical groups may be expanded to include dentists, psychologists, podiatrists, chiropractors, optometrists), mid-level providers (e.g., physician assistants, advanced practice nurses, nurse practitioners, certified nurse midwives) and allied-health professionals (e.g., certified nursing aides, medical assistants, radiology technicians, laboratory technicians, pharmacists, dietitians/nutritionists). Current sampling frames such as those from the American Medical Association may be obtained and studied, as well as frames that are not currently in use by NAMCS, such as state and organizational listings of other licensed providers; (2) within the National Study of Long-Term Care Providers, additional new frames may be sought and evaluated and data items from home care agencies, long-term care hospitals, and facilities exclusively serving individuals with intellectual/developmental disability may be tested. Similarly, data may be obtained from lists compiled by states and other organizations. Data about the facilities as well as residents and their visits will be investigated; and (3) in the inpatient and outpatient care settings, the National Hospital Care Survey (NHCS)

and the National Hospital Ambulatory Medical Care Survey (NHAMCS) may investigate the addition of facility and patient information especially as it relates to insurance and electronic medical records.

Projects under development or in the planning stages include two projects related to opioid use: one that will investigate adding questions to NAMCS on physician understanding of guidelines for opioid use and one that will test the validation of an algorithm for identifying opioid-involved hospital visits. Another study will develop a Hospital-Based Victim Services Frame.

The NHCSs collect critical, accurate data that are used to produce reliable national estimates—and in recent years (when budget allows), state-level estimates—of clinical services and of the providers who delivered those services in inpatient, outpatient, ambulatory, and long-term care settings. The data from these surveys are used by providers, policy makers and researchers to address important topics of interest, including the quality and disparities of care among populations, epidemiology of medical conditions, diffusion of technologies, effects of policies and practice guidelines, and changes in health care over time. Research studies need to be conducted to improve existing and proposed survey design and procedures of the NHCSs, as well as to evaluate alternative data collection approaches particularly due to the expansion of electronic health record use, and to develop new sample frames of currently out-of-scope providers and settings of care.

Average burdens are designed to cover 15–40 min interviews as well as 90-minute focus groups, as well as longer on-site visits, and situations where organizations may be preparing electronic data files. CDC requests OMB approval for an estimated 3,000 annual burden hours. There is no cost to respondents other than their time to participate.

TABLE 1—ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Health Care Providers and Business entities	Interviews, surveys, focus groups, experiments (in person, phone, internet, postal/electronic mail).	2,582	1	1
Health Care Providers, State/local government agencies, and business entities.	Interviews, surveys, focus groups, experiments (in person, phone, internet, postal/electronic mail).	167	1	2.5

Jeffrey M. Zirger,

Lead, Information Collection Review Office,  
Office of Scientific Integrity, Office of Science,  
Centers for Disease Control and Prevention.

[FR Doc. 2022-27504 Filed 12-19-22; 8:45 am]

BILLING CODE 4163-18-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-23-1185]

**Agency Forms Undergoing Paperwork Reduction Act Review**

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Youth Outreach Generic Clearance for the National Center for Health Statistics (NCHS)” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on September 23, 2022 to obtain comments from the public and affected agencies. CDC did not receive any comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street, NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

**Proposed Project**

Youth Outreach Generic Clearance for the National Center for Health Statistics (NCHS) (OMB Control No. 0920-1185, Exp. 07/31/2023)—Extension—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

The National Center for Health Statistics (NCHS) is authorized to collect data under Section 306 of the Public Health Service Act (42 U.S.C. 242k). NCHS has a history of reaching out to young people to encourage their interest in Science, Technology, Engineering and Math (STEM).

Examples of past involvement include adopting local schools, speaking at local colleges, conducting a Statistics Day for high school students, and, most recently, conducting the NCHS Data Detectives Camp for middle school students.

The success of these programs has inspired NCHS leadership and staff to want to look for new and continuing opportunities to positively impact the lives of young people and expand their interest, understanding of, and involvement in the sciences. NCHS requests approval for a New Generic Clearance mechanism to collect information for these youth outreach activities and to inform future NCHS planning activities.

As stated, these activities include hosting the Data Detectives Camp annually; hosting Statistics Day annually; creating youth poster sessions for professional conferences (such as the NCHS National Conference on Health Statistics or the American Statistical Association Conference etc.); hosting a statistical or health sciences fair or other STEM-related competitions; organizing a STEM Career Day or similar activity; developing web-based sites or materials with youth focus, as well as other programs developed to meet future youth outreach needs, particularly activities that encourage STEM.

Information will be collected using a combination of methodologies appropriate to each program. These may include: registration forms, letters of recommendation, evaluation forms, mail surveys, focus groups, automated and electronic technology (e.g. email, Web-based surveys), and telephone surveys.

OMB approval is requested for three years to conduct the Youth Outreach Generic Clearance for the National Center for Health Statistics (NCHS). CDC requests OMB approval for an estimated 1,750 annual burden hours. Participation is voluntary and there are no costs to respondents other than their time.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Type of survey	Respondent	Number of respondents	Number of responses/ respondent	Average burden/ response (in hours)
Questionnaires/Applications .....	Student/Youth .....	800	1	30/60
Applicants Questionnaire/Application .....	Parents/Guardians of Applicants .....	800	1	30/60
Applications, Recommendations, and Other applicant-supporting documentation.	School Officials/Community Representatives	1200	1	30/60
Focus Groups .....	Student/Youth; Parent/Guardian; School Officials; Other.	50	1	60/60
Other Program Surveys .....	Student/Youth; Parent/Guardian; School Officials; Other.	600	1	30/60