accredited. Accreditation is granted for a five-year period and the first several health departments have successfully completed the reaccreditation process. Formal efforts to assess the outcomes of the accreditation program began in late 2012 and continue to date. Priorities focus on gathering feedback for program improvement and documenting program outcomes to demonstrate impact and inform decision making about future program direction. Starting in 2012 and running through December 2019, the Robert Wood Johnson Foundation (RWJF) and the social science organization NORC at the University of

Chicago, led evaluation efforts. CDC will assume support of the evaluation starting in 2020 and as a result, OMB approval for data collection is being sought.

The purpose of this ICR is to support the collection of information from participating health departments through a series of five surveys. The surveys seek to collect longitudinal data on each health department throughout their accreditation process.

The respondent universe will include STLT health department directors or designees. All surveys will be administered electronically; a link to the survey website will be provided in the email invitation. The surveys will be administered on a quarterly basis and sent to all health departments that reach each milestone in the accreditation process (application, recently accredited, accredited for one year, approaching reaccreditation, and reaccreditation). Each health department will be invited to participate in each survey once (for a total of 5 surveys max per health department). The total annualized estimated burden is 100 hours. 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	Average burden per response (in hours)
STLT HD Directors or Designee	Survey 1: Applicants	60 60 60 60	1 1 1 1	20/60 20/60 20/60 20/60 20/60

Jeffrey M. Zirger,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-20-1097]

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 Monitoring and Reporting System for the National Tobacco Control Program 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 April 23, 2019 to obtain comments from the public and affected agencies. CDC received one comment 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

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov*. 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

Monitoring and Reporting System for the National Tobacco Control Program (0920–1097)—Reinstatement with Change—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) works with states, territories, tribal organizations, and the District of Columbia (collectively referred to as "state-based" programs) to develop, implement, manage, and evaluate tobacco prevention and control programs. Support and guidance for these programs have been provided through cooperative agreement funding and technical assistance administered by CDC's National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP). Partnerships and collaboration with other federal agencies, nongovernmental organizations, local communities, public and private sector organizations, and major voluntary associations have been critical to the success of these efforts. NCCDPHP cooperative agreements DP15-1509 (National State-Based Tobacco Control Programs) and DP14-1410PPHF14 (Public Health Approaches for Ensuring Quitline Capacity) continue to support efforts

since 1999 to build state health department infrastructure and capacity to implement comprehensive tobacco prevention and control programs. Through these cooperative agreements, health departments in all 50 states, the District of Columbia, Puerto Rico and Guam are funded to implement evidence-based environmental, policy, and systems strategies and activities designed to reduce tobacco use, secondhand smoke exposure, tobacco related disparities and associated disease, disability, and death. CDC plans to request OMB approval to collect information from the 53 statebased programs funded under both DP15–1509 and DP14–1410PPHF14. Awardees will report information about their work plan objectives, activities, infrastructure, and performance measures. Each awardee will submit an Annual Work Plan Progress Report using an Excel-based Work Plan Tool. The estimated burden per response on each of the abovementioned tools is six hours for each. Each awardee will submit an Annual Performance Measure report using an Excel-based Performance Measures tool. The

estimated burden per response on each of the abovementioned tools is five hours for each. Each awardee will submit an Annual Progress Report (APR) using an Excel-based APR tool. The estimated burden per response on each of the abovementioned tools is 18 hours for each. Each awardee will submit an Annual Component Model of Infrastructure (CMI) using an Excelbased CMI tool. The estimated burden per response on each of the abovementioned tools is three hours for each. In addition, each awardee will submit an Annual Budget Progress Report using an Excel-based Budget Tool. The estimated burden per response is five hours for each Annual Budget Progress Report. The same instruments will be used for all information collection and reporting throughout the OMB approval period. Awardees will upload their information to www.grantssolutions.gov on an annual basis to satisfy routine cooperative agreement reporting requirements.

CDC will use the information collected to monitor each awardee's progress and to identify facilitators and

challenges to program implementation and achievement of outcomes. Monitoring allows CDC to determine whether an awardee is meeting performance and budget goals and to make adjustments in the type and level of technical assistance provided to them, as needed, to support attainment of their performance measures. Monitoring and evaluation activities also allow CDC to provide oversight of the use of federal funds, and to identify and disseminate information about successful prevention and control strategies implemented by awardees. These functions are central to NCCDPHP's broad mission of reducing the burden of chronic diseases. Finally, the information collection will allow CDC to monitor the increased emphasis on partnerships and programmatic collaboration, and is expected to reduce duplication of effort, enhance program impact and maximize the use of federal funds.

OMB approval is requested for three years. Participation in the information collection is required as a condition of funding. 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	Average burden per response (in hours)
State Tobacco Control Managers	Annual Work Plan Progress Report	53 53 53 53 53	1 1 1 1	6 5 5 3 18

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention. [FR Doc. 2020–01048 Filed 1–22–20; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-20-0621; Docket No. CDC-2019-0117]

Proposed Data Collections Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the information collection project entitled National Youth Tobacco Surveys (NYTS) 2021–2023 which aims to collect data on tobacco use among middle- and high school students.

DATES: CDC must receive written comments on or before March 23, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2019-0117 by any of the following methods:

• Federal eRulemaking Portal: Regulations.gov. Follow the instructions for submitting comments. Mail: Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS-D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to Regulations.gov.

Please note: Submit all comments through the Federal eRulemaking portal (*regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office,