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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–20–0621]

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 National Youth Tobacco Survey 2021–2023 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 January 23, 2020 to obtain comments from the public and affected agencies. CDC received six 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.* 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

National Youth Tobacco Survey 2021–2023 (OMB Control No. 0920–0621, Exp. 4/30/2021)—Revision—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC)

Background and Brief Description

Tobacco use is the leading cause of preventable disease and death in the United States, and nearly all tobacco use begins during youth and young adulthood. A limited number of health risk behaviors, including tobacco use, account for the overwhelming majority of immediate and long-term sources of morbidity and mortality. Because many health risk behaviors are established during adolescence, there is a critical need for public health programs directed towards youth, and for information to support these programs.

Since 2004, the Centers for Disease Control and Prevention (CDC) has periodically collected information about tobacco use among adolescents (National Youth Tobacco Survey (NYTS) 2004, 2006, 2009, 2011, 2012, 2013–2020, OMB Control No. 0920–0621, Exp. 04/30/2021). This surveillance activity builds on previous surveys funded by the American Legacy Foundation in 1999, 2000, and 2002.

At present, the NYTS is the most comprehensive source of nationally representative tobacco data among students in grades 9–12. Moreover, the NYTS is the only source of such data for students in grades 6–8. The NYTS has provided national estimates of tobacco use behaviors, information about exposure to pro- and anti-tobacco influences, and information about racial and ethnic disparities in tobacco-related topics. Information collected through the NYTS is used to identify trends over time, to inform the development of tobacco cessation programs for youth, and to evaluate the effectiveness of existing interventions and programs.

CDC plans to request OMB approval to conduct additional cycles of the NYTS in 2021, 2022, and 2023. The survey will be conducted among nationally representative samples of students attending public and private schools in grades 6–12 and will be administered to students as a digitally-based survey programmed onto tablets. Information supporting the NYTS also will be collected from state-, district-, and school-level administrators and teachers. During the 2021–2023 timeframe, changes will be incorporated that reflect CDC's ongoing collaboration with FDA and the need to measure progress toward meeting strategic goals established by the Family Smoking Prevention and Tobacco Control Act. Information collection will occur annually and may include a number of new questions, as well as increased representation of minority youth.

The survey will examine the following topics: Use of e-cigarettes, cigarettes, cigars, smokeless tobacco, hookahs, roll-your own-cigarettes, pipes, snus, dissolvable tobacco, bidis, heated tobacco products, and nicotine pouches; knowledge and attitudes; media and advertising; access to tobacco products and enforcement of restrictions on access; secondhand smoke and e-cigarette aerosol exposure; provision of school- and community-based interventions, and cessation.

Results of the NYTS will continue to be used to inform and evaluate the National Comprehensive Tobacco Control Program; provide data to inform the Department of Health and Human Service's Tobacco Control Strategic Action Plan, and provide national benchmark data for state-level Youth Tobacco Surveys. Information collected through the NYTS also is expected to provide multiple measures and data for monitoring progress on seven tobacco-related objectives for Healthy People 2030.

OMB approval will be requested for three years. There are no costs to respondents other than their time. The total annualized burden is estimated to be 18,733 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr.)
State Administrators	State-level Recruitment Script for the NYTS	33	1	30/60
District Administrators	District-level Recruitment Script for the NYTS	253	1	30/60
School Administrators	School-level Recruitment Script for the NYTS	281	1	30/60
Teachers	Data Collection Checklist	1,177	1	15/60
Students	National Youth Tobacco Survey	24,000	1	45/60
	Cognitive Testing	40	1	120/60
	Survey Pre-tests	30	1	45/60
	Testing Activities	300	1	10/60

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–20–20AZ]

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 Evaluation of the Effectiveness of the Training and Education Modules in the North American Fatigue Management 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 November 4, 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 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* 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

Evaluation of the Effectiveness of the Training and Education Modules in the North American Fatigue Management Program—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC)

Background and Brief Description

The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety and health at work for all people through research and prevention. Reducing fatigue-related crashes is one of the top 10 changes needed to reduce transportation accidents and save lives identified by the National Transportation Safety Board (NTSB) and a National Occupational Research Agenda (NORA) priority. Fatigue is a preventable cause of crashes.

The North American Fatigue Management Program (NAFMP) was developed by the FMCSA, Transport Canada, and other entities to address commercial motor vehicle (CMV) driver fatigue through a comprehensive approach that delivers prevention information to carriers, dispatchers, drivers, and family members. In 2015, the National Academy of Sciences published the report ''Commercial motor vehicle driver fatigue, long-term health, and highway safety research needs'' that identified the need for fully evaluating the NAFMP so that recommendations for implementation of NAFMP are supported by scientific evidence. NIOSH is collaborating with the FMCSA to ensure the success of the proposed study.

NIOSH will recruit two commercial vehicle carriers, and CMV drivers, hereafter referred to as ''drivers'', employed by those carriers. Data will be collected during drivers' application to participate in the study, briefing session, study participation, and debriefing session. Data collection will primarily focus on driving performance, sleep, and sleepiness. These outcomes will be compared between pre-rollout of the NAFMP (in which drivers will operate as they did before their participation in the study) and after the rollout of the NAFMP training and education modules (in which drivers and managers will operate with increased knowledge, strategies, and techniques to reduce their fatigue). All drivers interested in participating in the study may complete the application. A briefing session will be scheduled with drivers who are found eligible for the study. During the briefing session, drivers who provide informed consent