

Thus Title X service sites see a large proportion of young and uninsured individuals. Over the past years, OPA has encouraged grantees to develop enrollment programs to ensure that clients who are currently uninsured understand new health insurance options that are available as a result of the ACA. Some sites already assist individuals with enrolling in Medicaid and other public insurance programs. With the availability of the health insurance marketplace, many more service delivery sites are assisting clients enroll in health insurance programs.

OPA does not have any data on how many sites are assisting and enrolling clients into health insurance programs. Thus we seek to collect this data in order to understand the impact of Title X funded service sites on assisting and enrolling clients into insurance programs. We will utilize this information to guide strategic planning around how Title X service sites and prepare for, and assist with, the full implementation of the ACA. Through a separate data collection process called the Family Planning Annual Report (FPAR) (OMB No. 0990–0221, expiration January 31, 2016), OPA collects information on the insurance

status of the clients served. With the implementation of the ACA, many of the traditional clients served by Title X service sites will qualify for health insurance. Due to the varying resources available at the State level to conduct outreach and enrollment, OPA has authorized grantees to use funding to conduct outreach and enrollment activities. However, we are not currently collecting data on how many sites are conducting such activities, the impact of those activities in enrolling clients into health insurance programs, and the need for additional resources to conduct outreach and enrollment. By collecting information on how many clients are assisted and enrolled in health insurance programs, OPA can; (1) measure the impact of Title X service sites in enrolling clients into insurance programs; (2) design strategic initiatives to encourage outreach and enrollment; and (3) better understand the impact of the Affordable Care Act on Title X service delivery sites.

Likely Respondents: This annual reporting requirement is for family planning services delivery projects authorized and funded by the Title X Family Planning Program [“Population Research and Voluntary Family Planning Programs” (Pub. L. 91–572)],

which was enacted in 1970 as Title X of the Public Health Service Act (Section 1001 of Title X of the Public Health Service Act, 42 United States Code [USC] 300).

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information.

This data is currently being collected by the Health Resources and Services Administration (HRSA) and the burden estimate is based on the supporting statement from their OMB application.

The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Outreach and Enrollment Activities	95	1	1	95

OS specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency’s functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Darius Taylor,
 Deputy, Information Collection Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60-Day–14–14BE]

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 404–639–7570 or send comments to CDC/ATSDR LeRoy Richardson, 1600 Clifton Road, MS D–

74, Atlanta, GA 30333 or send an email 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 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. Written comments should be received within 60 days of this notice.

Proposed Project

CDC Worksite Health Scorecard—New—National Center for Chronic Disease Prevention and Health

Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In the United States, chronic diseases such as heart disease, obesity and diabetes are among the leading causes of death and disability. Although chronic diseases are among the most common and costly health problems, they are also among the most preventable. Adopting healthy behaviors—such as eating nutritious foods, being physically active and avoiding tobacco use—can prevent the devastating effects and reduce the rates of these diseases.

Employers are recognizing the role they can play in creating healthy work environments and providing employees with opportunities to make healthy lifestyle choices. To support these efforts, the Centers for Disease Control and Prevention (CDC) plans to develop an online organizational assessment tool called the CDC Worksite Health Scorecard.

The CDC Worksite Health Scorecard is authorized by the Public Health Service Act and funded through the Prevention and Public Health Fund of the Patient Protection and Affordable Care Act (ACA). The CDC Worksite Health Scorecard is a tool designed to help employers assess whether they have implemented evidence-based health promotion interventions or strategies in their worksites to prevent heart disease, stroke, and related conditions such as hypertension, diabetes, and obesity. The assessment contains 125 yes/no questions that assess how evidence-based health promotion strategies are implemented at a worksite. These strategies include health promoting counseling services, environmental supports, policies, health plan benefits, and other worksite

programs shown to be effective in preventing heart disease, stroke, and related health conditions. Employers can use this tool to assess how a comprehensive health promotion and disease prevention program is offered to their employees, to help identify program gaps, and to prioritize across the following health topics: Organizational Supports; Tobacco Control; Nutrition; Physical Activity; Weight Management; Stress Management; Depression; High Blood Pressure; High Cholesterol; Diabetes; Signs and Symptoms of Heart Attack and Stroke; Emergency Response to Heart Attack and Stroke; Lactation Support; Community Resources; Occupational Health and Safety; and Vaccine-Preventable Diseases.

Employers, human resource managers, health benefit managers, health education staff, occupational nurses, medical directors, wellness directors, or others responsible for worksite health promotion can use the CDC Worksite Health Scorecard to establish benchmarks for their organizations and track improvements over time. State health departments may assist employers and business coalitions in using the tool and help them find ways to establish healthier workplaces. State health departments also can use the tool for monitoring worksite practices, establishing best practice benchmarks, and more effectively directing resources to support employers. Employers who complete the CDC Worksite Health Scorecard will be provided with workplace health program planning and implementation tools. Participating employers may also receive technical assistance and training.

The CDC Worksite Health Scorecard is a voluntary, self-reported online

survey that will be available to any public/private employer regardless of size, industry sector, or geographic location. The online system will require the creation of a user account with employer contact information so that employer representatives can complete the CDC Worksite Health Scorecard instrument; receive an immediate feedback report on existing program gaps; and benchmark themselves against other employers using the CDC Worksite Health Scorecard. It is recommended that the CDC Worksite Health Scorecard be repeated on an annual basis.

CDC is requesting Office of Management and Budget (OMB) approval by March 2014. The information to be collected will allow CDC to register employers and permit access to the survey and other resources such as the user manual, feedback reports, and tools for employers. CDC will also use the information to generate benchmark reports for comparing the number of workplace health strategies an individual employer has implemented to the number of strategies implemented by other employers using the CDC Worksite Health Scorecard, to identify success drivers for building and maintaining successful workplace health programs, to raise awareness and knowledge among employers about science-based workplace health program strategies, to develop additional tools and resources for employers, and to evaluate the impact of the CDC Worksite Health Scorecard on the adoption of workplace health programs, policies and environmental supports.

OMB approval is requested for three years. CDC estimates that 600 employers will complete the CDC Worksite Health Scorecard per year. Participation is voluntary and there are no costs to participants 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 hr)	Total burden (in hr)
Employers	CDC Worksite Health Scorecard	600	1	30/60	300
Total	300

LeRoy Richardson,
*Chief, Information Collection Review Office,
 Office of Scientific Integrity, Office of the
 Associate Director for Science, Office of the
 Director, Centers for Disease Control and
 Prevention.*

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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

**Centers for Disease Control and
 Prevention**

[60-Day-14-0210]

**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 404-639-7570 or send comments to CDC, LeRoy Richardson, 1600 Clifton Road, MS D-74, Atlanta, GA 30333 or send an email 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 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. Written comments should be received within 60 days of this notice.

Proposed Project

List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products (OMB No. 0920-0210, exp. 2/28/2014)—Extension—Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Cigarette smoking is the leading preventable cause of premature death and disability in the United States. Each year, more than 443,000 premature deaths occur as the result of diseases related to cigarette smoking. The Centers for Disease Control and Prevention (CDC), Office on Smoking and Health (OSH) has the primary responsibility for the Department of Health and Human Services (HHS) smoking and health program. HHS's overall goal is to reduce death and disability resulting from cigarette smoking and other forms of tobacco use through programs of information, education and research.

The Comprehensive Smoking Education Act of 1984 (CSEA, 15 U.S.C. 1336 or Pub. L. 98-474) requires each person who manufactures, packages, or imports cigarettes to provide the Secretary of HHS with a list of ingredients added to tobacco in the manufacture of cigarettes. The legislation also authorizes HHS to undertake research, and to report to the Congress (as deemed appropriate) discussing the health effects of these ingredients.

HHS has delegated responsibility for implementing the CSEA's ingredient reporting requirements to CDC's OSH. OSH has collected ingredient reports on cigarette products since 1986.

Respondents are commercial cigarette manufacturers, packagers, or importers, or their designated representatives. Respondents are not required to submit specific forms; however, they are required to submit a list of all ingredients used in their products. CDC requires the ingredient report to be submitted by chemical name and Chemical Abstract Service (CAS) Registration Number, consistent with accepted reporting practices for other companies currently required to report ingredients added to other consumer products. Typically, respondents submit a summary report to CDC with the ingredient information for multiple products, or a statement that there are no changes to their previously submitted ingredient report. The estimated burden per response is 6.5 hours. The total estimated annualized burden hours are 501.

Ingredient reports for new products are due at the time of first importation. Thereafter, ingredient reports are due annually on March 31. Information is submitted to OSH by mailing a written report on the respondent's letterhead, which may be accompanied by a compact disk (CD), three-inch floppy disk, or thumb drive. Annual ingredient reports should be mailed to: Office on Smoking and Health, Attention: FCLAA Program Manager, National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention, 4770 Buford Highway, NE., MS F-79 Atlanta, GA 30341-3717. Electronic mail submissions are not accepted. Upon receipt and verification of the annual ingredient report, OSH issues a Certificate of Compliance to the respondent.

There are no costs to respondents other than their time. Office of Management and Budget (OMB) approval is requested for three years.

Estimated Annualized Burden Hours

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Cigarette Manufacturers, Packagers, and Importers	77	1	6.5	501