

Leroy A. 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.*
[FR Doc. 2013–25860 Filed 10–30–13; 8:45 am]
BILLING CODE 4163–18–P

**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**
[60Day–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

Leroy A. 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.

[FR Doc. 2013-25799 Filed 10-30-13; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-14-0879]

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 LeRoy Richardson, 1600 Clifton Road, MS-D74, 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

Surveys of State, Tribal, Local, and Territorial (STLT) Governmental Agencies (OMB Control No. 0920-0879, Exp. 3/31/2013)—Revision—Office of the Director, Office for State, Tribal Local and Territorial Support (OSTLTS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC's mission is to create the expertise, information, and tools that people and communities need to protect their health—through health promotion, prevention of disease, injury and disability, and preparedness for new health threats. CDC seeks to accomplish its mission by collaborating with partners throughout the nation and the world to: Monitor health, detect and investigate health problems, conduct research to enhance prevention, develop and advocate sound public health policies, implement prevention strategies, promote healthy behaviors, foster safe and healthful environments, and provide leadership and training.

CDC is requesting a three-year approval for a generic clearance to collect information related to domestic public health issues and services that

affect and/or involve state, tribal, local and territorial (STLT) government entities. The respondent universe is comprised of STLT governmental staff or delegates acting on behalf of a STLT agency involved in the provision of essential public health services in the United States. Delegate is defined as a governmental or non-governmental agent (agency, function, office or individual) acting for a principal or submitted by another to represent or act on their behalf. The STLT agency is represented by a STLT entity or delegate with a task to protect and/or improve the public's health. Information will be used to assess situational awareness of current public health emergencies; make decisions that affect planning, response and recovery activities of subsequent emergencies; fill CDC gaps in knowledge of programs and/or STLT governments that will strengthen surveillance, epidemiology, and laboratory science; improve CDC's support and technical assistance to states and communities. CDC will conduct brief data collections, across a range of public health topics related to essential public health services.

CDC estimates up to 30 data collections with STLT governmental staff or delegates, and 10 data collections with local/county/city governmental staff or delegates will be conducted on an annual basis. Ninety-five percent of these data collections will be web-based and five percent telephone, in-person, and focus groups. The total annualized burden of 54,000 hours is based on the following estimates.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of surveys per respondent type	Average burden per respondent (in hours)	Total burden hours (annual)
State, Territorial, or Tribal government staff or delegate.	Web, telephone, in-person, focus group.	800	30	1	24,000
Local/County/City government staff or delegate.	Web, telephone, in-person, focus group.	3,000	10	1	30,000
Total	54,000