DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–14–0210]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call (404) 639–7570 or send an email to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 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

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.

Since 1986, as required by the Comprehensive Smoking Education Act of 1984 (CSEA, 15 U.S.C. 1336 or Pub. L. 98–474), CDC has collected information about the ingredients used in cigarette products. Respondents are commercial cigarette manufacturers, packagers, or importers (or their representatives), who are required by the CSEA to submit ingredient reports to HHS on an annual basis.

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.

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, by CD, three-inch floppy disk, or thumb drive. Electronic mail submissions are not accepted. Mail Annual Ingredient Submissions to Attention: FCLAA Program Manager, Centers for Disease Control and Prevention, National Center for Chronic Disease Prevention and Health Promotion, Office on Smoking and Health, 4770 Buford Highway, NE., MS F–79, Atlanta, GA 30341–3717.

Upon receipt and verification of the annual ingredient report, OSH issues a Certificate of Compliance to the respondent. OSH also uses the information to report to the Congress (as deemed appropriate) discussing the health effects of these ingredients.

There are no costs to respondents other than their time. The annualized number of respondents is 77 and the total estimated annualized burden hours are 501. OMB approval is requested for three years.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Cigarette Manufacturers, Packagers, and Importers	Ingredient Report	77	1	6.5

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. 2014–01648 Filed 1–28–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–14–0338]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call (404) 639–7570 or send an email to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Annual Submission of the Ingredients Added to, and the Quantity of Nicotine Contained in, Smokeless Tobacco Manufactured, Imported, or Packaged in the U.S. (OMB No. 0920–0338, exp. 2/28/2014)—Extension—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), 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 the use of smokeless tobacco products and other forms of tobacco use through programs of information, education and research.

Since 1994, as required by the Comprehensive Smokeless Tobacco Education Act of 1986 (CSTHEA, 15 U.S.C. 4401 *et seq.,* Pub. L. 99–252), CDC has collected information about the ingredients used in smokeless tobacco products and their nicotine content. Respondents are commercial smokeless tobacco product manufacturers, packagers, or importers (or their designated representatives), who are

required by the CSTHEA to submit ingredient reports to HHS on an annual basis. The legislation also authorizes HHS to undertake research, and to report to Congress, as deemed appropriate, about the health effects of these ingredients.

Respondents are not required to submit specific forms; however, they are required to meet reporting guidelines and to submit the ingredient report 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.

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, by CD, three-inch floppy disk, or thumb drive. The information collection is subject to strict confidentiality provisions and electronic mail submissions are not accepted. Upon receipt and verification of the annual nicotine and ingredient report, OSH issues a Certificate of Compliance to the respondent.

OMB approval is requested for three years. There are no changes to information collection procedures, the estimated burden per response, or the estimated number of respondents. The total estimated annualized burden hours are 22,269. 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)
Smokeless Tobacco Manufacturers, Packagers, and Importers.	SLT Nicotine and Ingredient and Report	13	1	1,713

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. 2014–01649 Filed 1–28–14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2014–0003]

Draft Guideline—Centers for Disease Control and Prevention Draft Guideline for the Prevention of Surgical Site Infections

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

ACTION: Notice of availability and request for public comment.

SUMMARY: The Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS) requests public comment on the Draft Guideline for the Prevention of Surgical Site Infections (SSIs) (draft Guideline). The Draft Guideline addresses new and updated strategies for the prevention of SSI in healthcare settings. This draft Guideline can be found at *http://www.regulations.gov* Docket No. CDC–2014–0003. CDC is also publishing the supporting appendices that include primary evidence, study evaluation, and data evaluation tables that were used in developing the draft Guideline recommendations at *http://www.regulations.gov.*

The draft Guideline is designed for use by infection prevention staff, healthcare epidemiologists, administrators, nurses, and personnel responsible for developing, implementing, and evaluating infection prevention and control programs for healthcare settings across the continuum of care. The recommendations contained in the draft Guideline are based on a targeted systematic review of the best available evidence for specific topics related to the prevention of surgical site infections (SSI).

DATES: Comments must be received on or before February 28, 2014.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2014–0003, by any of the following methods:

- Federal eRulemaking Portal: *http://www.regulations.gov.* Follow the instructions for submitting comments.
- Mail: Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention, Attn: Guideline for the Prevention of Surgical Site Infections, Docket No. CDC–2014–0003, 1600 Clifton Road NE., Mailstop A07, Atlanta, Georgia 30333.

Instructions: All submissions received must include the agency name and docket number or RIN. All relevant public comments received will be posted publicly to *www.regulations.gov* without change, including any personal or proprietary information provided. To download an electronic version of the draft Guideline and appendices, access *http://www.regulations.gov.*

Written materials identified by Docket No. CDC–2014–0003 will be available during the comment period for public inspection Monday through Friday, except for legal holidays, 9 a.m. until 4:30 p.m. Eastern Standard Time, at CDC Library, 1600 Clifton Road NE., Atlanta, Georgia 30333. Please call ahead to (404) 639–1717 and request a Library representative schedule your visit. All public comments will be reviewed and considered prior to finalizing the draft Guideline.

FOR FURTHER INFORMATION CONTACT: Erin Stone, Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention, 1600 Clifton Road NE., Mailstop A–31, Atlanta, Georgia 30333; Telephone: (404) 639–4000.

SUPPLEMENTARY INFORMATION: Since 2010 CDC has collaborated with national partners, academicians, public and private health professionals, and other partners to create this draft Guideline. Additionally, CDC sought input in each phase of development from subject matter experts in surgery, infectious diseases, and orthopedics through a Guideline Expert Panel formed to develop the new draft Guideline. CDC also received input from the Healthcare Infection Control Practices Advisory Committee (HICPAC) throughout the development of the draft Guideline. HICPAC includes representatives from public health,