applied for a one-year extension that ends September 30, 2014.

CDC currently collects information from awardees about the activities supported with Traditional Foods funding. Twice per year, each awardee submits a shared data elements (SDE) report to CDC through a Web-based interface. The SDE are organized in three domains: Traditional Local Healthy Foods, Physical Activity, and Social Support for Healthy Lifestyle Change and Maintenance. Reports are submitted to CDC in the spring and fall. The spring 2014 report will be submitted to CDC under the current OMB clearance (OMB No. 0920–0889, exp. 6/30/2014).

CDC plans to request OMB approval of a six-month extension of the Traditional Foods information collection, through approximately December 31, 2014. The extension will enable CDC to receive a final report on activities conducted during late spring, summer, and early fall of 2014. Because of the variety of food- and lifestyle-related programs that take place in these seasons, CDC wants to ensure complete and accurate reporting of awardee activities conducted the last 5–6 months of cooperative agreement funding.

There are no proposed changes to the data collection instrument, data collection methodology, or the estimated burden per response. Changes to be implemented in this Revision request include: (1) A reduction in the number of respondents, from 17 to 16, (2) a change in the frequency of reporting (only one SDE report will be received during the six-month extension period), and (3) discontinuation of the one-time retrospective data collection that was part of the initial three-year clearance request.

CDC will continue to use the SDE reports to compile a systematic, quantifiable inventory of activities, products, and outcomes associated with the Traditional Foods program. The SDE also allow CDC to analyze aggregate data for improved technical assistance and overall program improvement, reporting, and identification of outcomes; allow CDC and grantees to create a comprehensive inventory/resource library of diabetes primary prevention ideas and approaches for AI/AN communities and identify emerging best practices; and improve dissemination of success stories. The SDE supplements the narrative progress reports that grantees submit to CDC in conjunction with the annual continuation application for funding. Although these reports provide important contextual information and are useful for local program monitoring, they do not support the production of statistical reports that are needed to fully describe the Traditional Foods program and to respond to various administrative inquiries.

Respondents will be 16 Tribes and Tribal organizations that receive funding through the Traditional Foods program. The SDE will continue to be submitted to CDC using Survey Monkey, an electronic Web-based interface. The estimated burden per response is two hours. Each grantee will receive a personalized advance notification letter, followed by an email with a link to the Survey Monkey site.

Participation in this information collection is required for Traditional Foods program awardees. 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 hrs)	Total burden (in hrs)
AI/AN Tribal Grantees	Traditional Foods Shared Data Elements.	16	1	2	32

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. 2014–02510 Filed 2–5–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention (CDC)

[CDC–2013–0024, Docket Number NIOSH–270]

NIOSH Center for Motor Vehicle Safety: Research and Guidance Strategic Plan 2014–2018

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of draft document for public comment.

SUMMARY: The National Institute for Occupational Safety and Health of the Centers for Disease Control and Prevention announces the availability of a draft document entitled *NIOSH Center for Motor Vehicle Safety: Research and Guidance Strategic Plan 2014–2018* for public comment. To view the notice and related materials, visit *http://www.regulations.gov* and enter CDC–2013–0024 in the search field and click ''Search.''

Public comment period: Comments must be received within 30 days from publication of the **Federal Register** Notice.

ADDRESSES: You may submit comments, identified by CDC–2013–0024 and Docket Number NIOSH–270, by either of the following two methods:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* NIOSH Docket Office, Robert A. Taft Laboratories, 4676 Columbia Parkway, MS C–34, Cincinnati, Ohio 45226.

Instructions: All information received in response to this notice must include the agency name and docket number (CDC–2013–0024; NIOSH–270). All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. All electronic comments should be formatted as Microsoft Word. Please make reference to CDC–2013–0024 and Docket Number NIOSH–270.

SUPPLEMENTARY INFORMATION: The purpose of this review is to receive public comments and input on the NIOSH Center for Motor Vehicle Safety: Research and Guidance Strategic Plan for the period 2014–2018. NIOSH is seeking comments on: (1) The relevance of the current draft; (2) the adequacy of the plan in addressing research needs for work-related motor vehicle crashes and fatal/non-fatal injuries; (3) the

adequacy of proposed performance measures; and (4) additional potential partners the NIOSH Center for Motor Vehicle Safety could engage with to enhance the relevance and capacity of the Center's program.

Background: Fatality data show that across all industries, motor vehicle-related incidents are consistently the leading cause of work-related fatalities, and they are the first or second leading cause in every major industry sector. The NIOSH Center for Motor Vehicle Safety is the focal point for research and prevention activities within the Institute to reduce work-related motor vehicle crashes and resulting injuries. The goals for the NIOSH Center for Motor Vehicle Safety were developed based on: (1) Consideration of research gaps based on review of the scientific literature, employer policies, and government regulations; (2) a review of related goals in the NIOSH sector and cross-sector programs; and (3) consideration of the research areas where NIOSH is best-positioned to add to the knowledge base on work-related motor vehicle safety. The draft goals address the following areas:

(1) Epidemiologic research to identify risk factors associated with work-related motor vehicle crashes and injury

(2) Engineering and technology-related research

(3) Research and demonstration projects on motor vehicle safety management strategies

(4) Global collaborations to develop strategies for reducing occupational road traffic injuries worldwide

(5) Research communication products

FOR FURTHER INFORMATION CONTACT: Stephanie Pratt, Ph.D., NIOSH, Division of Safety Research, Mailstop H–1808, 1095 Willowdale Road, Morgantown, West Virginia 26505–2888. Dr. Pratt may be contacted at (304) 285–5992 or by email at *sgp2@cdc.gov*.

Dated: January 31, 2014.

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2014–02524 Filed 2–5–14; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0623]

Agency Information Collection Activities; Proposed Collection; Comment Request; Voluntary Cosmetic Registration Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice invites comments on the collection of information associated with our Voluntary Cosmetic Registration Program (VCRP).

DATES: Submit either electronic or written comments on the collection of information by April 7, 2014.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov.* Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Voluntary Cosmetic Registration Program—21 CFR Parts 710 and 720 (OMB Control Number 0910–0027)—Extension

The Federal Food, Drug, and Cosmetic Act (the FD&C Act) provides us with the authority to regulate cosmetic products in the United States. Cosmetic products that are adulterated under section 601 of the FD&C Act (21 U.S.C. 361) or misbranded under section 602 of the FD&C Act (21 U.S.C. 362) may not be distributed in interstate commerce. We have developed the VCRP to assist us in carrying out our responsibility to regulate cosmetics.

In 21 CFR part 710, we request that establishments that manufacture or package cosmetic products register with us on Form FDA 2511 entitled ''Registration of Cosmetic Product Establishment.'' The term ''Form FDA 2511'' refers to both the paper and electronic versions of the form. The electronic version of Form FDA 2511 is available on our VCRP Web site at *http://www.fda.gov/Cosmetics/GuidanceComplianceRegulatoryInformation/VoluntaryCosmeticsRegistrationProgramVCRP/OnlineRegistration/default.htm.* We strongly encourage electronic registration of Form FDA 2511 because it is faster and more convenient. A registering facility will receive confirmation of electronic registration, including a registration number, by email, usually within 7 business days. The online system also