Dated: September 9, 2009.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E9–22141 Filed 9–14–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0487]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Safety Survey

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by October 15, 2009.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0345. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794, *JonnaLynn.Capezzuto@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Food Safety Survey—(OMB Control Number 0910–0345—Reinstatement)

Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(b)(2)), FDA is authorized to conduct research relating to foods and to conduct educational and public information programs relating to the safety of the nation's food supply. The Food Safety Survey is a nationally representative survey of consumers' knowledge, attitudes, and beliefs about food safety. Previous versions of the survey were collected in 1988, 1993, 1998, 2001, and 2006. Data from the previous surveys are being used to evaluate two Healthy People 2010 objectives: (1) Increase the proportion of consumers who follow key food safety practices (Objective 10–5), and (2) reduce severe allergic reactions to food among adults (Objective 10–4b). Additionally, data are used to measure trends in consumer food safety habits including hand and cutting board washing, cooking practices, and use of food thermometers. Finally, data are used to evaluate educational messages and to inform policymakers about consumer attitudes about novel technologies such as food irradiation and biotechnology.

Since 2006, there have been several high profile recalls of FDA-regulated food due to contamination. Information about food recalls does not always reach the intended audience (Refs. 1, 2, and 3). The Food Safety Survey planned for 2009 will look specifically at reasons why consumers do not always heed food recall alerts. A new food recall module will be added that contains new questions to learn about how recent food recalls have affected consumer confidence in the food supply and what effect, if any, they have on consumers' home food safety behaviors. This information will help FDA develop strategies to more effectively communicate food recall information to the public.

The methods for the 2009 version of the Food Safety Survey will be the same as for the previous Food Safety Surveys. A nationally representative sample of 4,000 adults in households with telephones will be selected at random and interviewed by telephone. This survey will include an oversample of Hispanics with a minimum of 500 Hispanics sampled. Additionally, 200 initial nonrespondents will be asked to participate in a short version of the survey to conduct a nonresponse analysis. Participation will be voluntary. Cognitive interviews and a pretest will be conducted prior to fielding the survey.

In the **Federal Register** of September 17, 2008 (73 FR 53878), FDA published a 60-day notice requesting public comment on the proposed collection of information. The agency received one comment that was not responsive to the comment request on the information collection provisions.

FDA estimates the burden of this collection of information as follows:

The total estimated burden imposed by this collection of information is 1,541 hours (table 1 of this document).

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Activity	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Cognitive Interview	20	1	20	1	20
Pretest	27	1	27	0.5	14
Screener	10,000	1	10,000	.0167	167
Survey	4,000	1	4,000	.33	1,320
Nonresponse	200	1	200	.10	20
Total					1,541

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Prior to finalizing the survey, FDA will conduct 20 cognitive interviews each requiring an average of 1 hour per respondent for a total of 20 hours. Before the survey is fielded, a small pretest of 27 individuals, each lasting half an hour (0.5 hour), will be conducted. The survey screener is estimated to take 1 minute or less per

response for a total screener burden of 4,000 (respondents) + 6,000 (ineligibles screened) x .0167 hours = 167 hours. The survey will require an average of 20 minutes (0.33 hours) per respondent and we expect that the variation in burden across respondents will be small. This estimate is based on average interview time for the 2006 Food Safety Survey. The proposed number of respondents is 4,000, each of whom will be asked to complete a one-time telephone interview that requires no preparation time. Additionally, 200 initial nonrespondents will be asked to participate in a short version of the survey to conduct a nonresponse analysis. This is expected to take 6 minutes (0.10 hours). Therefore, the total estimated public reporting burden is 1,541 hours.

We have revised the burden table. In the 60-day notice published on September 17, 2008, we estimated the total burden to be 1,421 hours. The total burden of 1,541 hours estimated in table 1 of this document includes an additional 120 hours, which resulted from correcting a typographical error in line 4 of the table. The hours per response in line 4 of table 1 changed from 0.3 to 0.33.

Dated: September 1, 2009.

David Horowitz,

Assistant Commissioner for Policy.

[FR Doc. E9–22121 Filed 9–14–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0406]

Agency Emergency Processing Under the Office of Management and Budget Review; Tobacco Product Establishment Registration and Submission of Certain Health Information; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of September 1, 2009 (74 FR 45219). The document announced the proposed collection of information concerning the submission of tobacco product establishment registration and submission of certain health information, including ingredient listing and health related documents, as required by the Family Smoking Prevention and Tobacco Control Act. The document was published with an incorrect date for submitting written or electronic comments on the proposed collection. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794, *Jonnalynn.Capezzuto@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In FR Doc. E9–21099, appearing on page 45219, in the **Federal Register** of Tuesday, September 1, 2009, the following correction is made:

On page 45219, in the second column, in the ''**DATES**'' section, beginning in the second line, ''September 16, 2009'' is corrected to read ''October 1, 2009''.

Dated: September 8, 2009.

David Horowitz,

Assistant Commissioner for Policy.

[FR Doc. E9–22120 Filed 9–14–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Injury Prevention and Control Initial Review Group: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the National Center for Injury Prevention and Control Initial Review Group, Department of Health and Human Services, has been renewed for a 2-year period through August 20, 2011.

For information, contact Dr. Richard Waxweiler, Executive Secretary, National Center for Injury Prevention and Control Initial Review Group, Department of Health and Human Services, 1600 Clifton Road, M/S F63, Atlanta, Georgia 30341, telephone 770/488–4850, or fax 770/488–4422.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and othercommittee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: September 4, 2009.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E9–22140 Filed 9–14–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, Public Health Service, HHS.

ACTION: Notice.

SUMMARY: The inventions listed below are owned by an agency of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 207 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

ADDRESSES: Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852–3804; telephone: 301/496–7057; fax: 301/402–0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

Purified Saxitoxin for Food Safety Applications

Description of Technology: Available for licensing as a biological material for research purposes is purified saxitoxin. Saxitoxin is the parent compound in a family of natural toxins that can occur in seafood and can cause food borne illness. Highly purified saxitoxin is vital for the development, validation, and calibration of detection methods for these toxins, as well as for fundamental studies in physiology and pain management. Interested parties may license the compound for conjugation chemistry and radiolabeling with the end goal of generating a research reagent.