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.