ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Type of form	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Program participant	Follow-up Survey Screen for Matter of Balance-Introduction Script.	500	1	3/60
	Follow-up Survey for Matter of Balance	425	1	45/60
Program coordinator	Cost assessment of AoA-funded fall prevention programs.	200	1	2

Dated: February 26, 2009.

Maryam Daneshvar,

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

[FR Doc. E9–4728 Filed 3–4–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Injury Prevention and Control Initial Review Group (NCIPC IRG)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), CDC announces the following meeting of the aforementioned review group:

Times and Dates:

6 p.m.–6:30 p.m., March 23, 2009 (Open).

6:30 p.m.–8 p.m., March 23, 2009 (Closed).

8 a.m.–5 p.m., March 24, 2009 (Closed).

8 a.m.–5 p.m., March 25, 2009 (Closed).

Place: The W Hotel, 3377 Peachtree Road, NE., Atlanta, Georgia 30326, Telephone: (678) 500–3181.

Status: Portions of the meetings will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5, U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Section 10(d) of Public Law 92–463.

Purpose: This group is charged with providing advice and guidance to the Secretary, Department of Health and Human Services, and the Director, CDC, concerning the scientific and technical merit of grant and cooperative agreement applications received from academic institutions and other public and private profit and nonprofit organizations, including State and local government agencies, to conduct specific injury research that focuses on prevention and control.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of applications submitted in response to Fiscal Year 2009 Requests for Applications related to the following individual research announcement: CE09–007, Research Grants for Preventing Violence and Violence Related Injury (R01).

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: J Felix Rogers, Phd, M.P.H., NCIPC, Extramural Research Program Office, CDC, 4770 Buford Highway, NE., M/S F62, Atlanta, Georgia 30341–3724, Telephone (770) 488–4334.

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

Dated: February 25, 2009.

Elaine L. Baker,

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

[FR Doc. E9–4642 Filed 3–4–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Subcommittee for Dose Reconstruction Reviews (SDRR), Advisory Board on Radiation and Worker Health (ABRWH), National Institute for Occupational Safety and Health (NIOSH)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention, announces the following meeting for the aforementioned subcommittee:

Time and Date:

9:30 a.m.–5 p.m., March 12, 2009.

Place: Cincinnati Airport Marriott, 2395 Progress Drive, Hebron, Kentucky 41018. Telephone (859) 334–4611, Fax (859) 334–4619.

Status: Open to the public, but without a public oral comment period. To access by conference call dial the following information 1 (866) 659–0537, Participant Pass Code 9933701.

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines that have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC).

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2009.

Purpose: The Advisory Board is charged with (a) Providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee for Dose Reconstruction

Reviews was established to aid the Advisory Board in carrying out its duty to advise the Secretary, HHS, on dose reconstruction.

Matters To Be Discussed: The agenda for the Subcommittee meeting includes: the selection of an 11th set of dose reconstructions for review; discussion of cases under review from the 6th, 7th, and 8th sets of individual dose reconstructions; preparation of a letter report on the first 100 dose reconstruction cases reviewed; and, discussion of selection criteria and review rate for 2009.

The agenda is subject to change as priorities dictate. Written comments may be submitted from the public. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

This meeting was previously scheduled to convene on January 29, 2009, but was cancelled due to inclement weather and airport facility inaccessibility. The meeting was scheduled to reconvene as soon as possible; therefore, this **Federal Register** notice is being published less than fifteen days prior to the meeting date.

Contact Person for More Information: Theodore Katz, Designated Federal Officer, CDC, NIOSH, 1600 Clifton Road, Mailstop E–20, Atlanta, GA 30333, Telephone (513) 533–6800, Toll Free 1 (800) CDC–INFO, E-mail *ocas@cdc.gov.*

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

Dated: February 26, 2009.

Elaine L. Baker,

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

[FR Doc. E9–4643 Filed 3–4–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0553]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Survey To Evaluate the Effectiveness of Mississippi Delta Fish Advisories

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 April 6, 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–NEW and title ''Survey to Evaluate the Effectiveness of Mississippi Delta Fish Advisories.'' 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.

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.

Survey to Evaluate the Effectiveness of Mississippi Delta Fish Advisories—(OMB Control Number 0910–NEW)

The proposed survey will gather information about fishing and fish consumption habits in the Mississippi Delta region, as well as the respondents' awareness and understanding of the Regional Delta Advisory (RDA) issued by the Mississippi Department of Environmental Quality. 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. In June 2005, the Environmental Protection Agency's (EPA's) Office of Water and FDA's Center for Food Safety and Applied Nutrition finalized a Memorandum of Understanding (MOU) to enhance collaboration between FDA and EPA regarding environmental contaminants in fish and shellfish and the safety of fish and shellfish for U.S. consumers. The MOU is available at *http://www.epa.gov/waterscience/fish/files/moufdaepa.pdf*.

The proposed study is phase two of a two phase study designed to determine whether existing fish consumption recommendations issued by the State of Mississippi are adequately protecting sport and subsistence consumers of fish harvested from Delta waters. The final report of phase one, entitled ''Recommended Study Design for a Survey to Evaluate the Effectiveness of Mississippi Delta Fish Advisories,'' is available at *http://www.epa.gov/waterscience/fish/technical/ms-delta.html*. Based on the report cited in this paragraph, FDA is conducting the proposed survey on behalf of EPA to evaluate the effectiveness of the Mississippi Delta Fish Advisories. The proposed survey will collect information on the extent to which Delta sport and subsistence fishermen and their families are aware of the RDA and its recommendations and the extent to which the respondents have changed their fish consumption behaviors as a result of the advisory. The survey will also document specific behavior changes resulting from the RDA, such as increases or decreases in the amount of locally harvested fish consumed, changes in methods of fish preparation, and consumption or avoidance of specific species of fish.

Results of the survey will provide EPA information about fishing and fish consumption habits in the Mississippi Delta region, as well as the respondents' awareness and understanding of the RDA.

The respondents will be selected from four counties in the Mississippi Delta region. Counties were selected to include a mix of rural and non-rural areas and areas with major water resources affected by the advisory. The selected counties are Coahoma, Holmes, Leflore, and Washington. Only the part of Holmes County that is within the advisory area will be included in the survey.

The total sample will include 400 on-the-banks interviews and 600 household interviews of sport and subsistence fishers who harvest noncommercial fish