Dated: December 23, 2010.

David Dorsey,

Acting Deputy Commissioner for Policy, Planning and Budget.

[FR Doc. 2010–32948 Filed 12–29–10; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0118]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 28, 2010 (75 FR 30036), the Agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0520. The approval expires on December 31, 2013. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: December 23, 2010.

David Dorsey,

Acting Deputy Commissioner for Policy, Planning and Budget.

[FR Doc. 2010–32946 Filed 12–29–10; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Testing Successful Health Communications Surrounding Aging-Related Issues From the National Institute on Aging (NIA)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute on Aging, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on 09–27–2010 at 08:45:00 (*http://federalregister.gov/a/2010–24277,* Vol. 75, No. 187, Page: 59723–59724 (2 pages); Document Citation: 75 FR 59723; Document Number: 2010–24277) and allowed 60-days for public comment. One comment was received from an organization who requested to be considered as a contractor for NIA's project. No other public comments were received. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: *Title:* Testing successful health communications surrounding aging-related issues from the National Institute on Aging (NIA*). Type of Information Collection Request:* New. *Need and Use of Information Collection:* This study will support NIA's mission "to communicate information about aging and advances in research on aging to the scientific community, health care providers, and the public." The primary objectives of this study are to:

- Assess audiences' trusted/preferred sources for information, knowledge, attitudes, behaviors, and other characteristics for the planning/development of health messages and communications strategies;
- Pre-test health messages and outreach strategies while they are in developmental form to assess audience response, including their likes and dislikes.

NIA's Office of Communications and Public liaison will collect this information through formative qualitative research with its key audiences—older people, caregivers, and health professionals. Methods will include focus groups, individual interviews, self-administered questionnaires, and website surveys. The information will be used to (1) develop and revise health information resources and outreach strategies to maximize their effectiveness; (2) determine new topic areas to explore for future NIA publications; and (3) identify new ways to support the health information needs of older adults and people who serve older adults. NIA is requesting a generic clearance for a range of research data collection procedures to ensure that they successfully develop and disseminate effective health communications on aging-related issues. *Frequency of Response:* On occasion. *Affected Public:* Older people, caregivers, and health professionals (physicians and non-physicians). *Type of Respondents:* Older people, caregivers, and health professionals (physicians and non-physicians). The annual reporting burden is as follows: *Estimated Number of Respondents:* 630. *Estimated Number of Responses per Respondent:* 1. *Average Burden Hours per Response:* 0.37. *Estimated Total Annual Burden Hours Requested:* 234. The annualized cost to respondents is estimated at: $5,680. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Type of respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total annual burden hours requested
Older adults	260	1	.37	97
Non-physician health professionals and caregivers	310	1	.35	107
Physicians	60	1	.5	30
Total				234

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, *Attention:* Desk Officer for NIH. *To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact:* Megan Homer, Writer/Editor, Office of Communications and Public Liaison, NIH, Building 31C Room 5C27, 9000 Rockville Pike, Bethesda, MD 20892, or call non-toll-free number 301–496–1752 or E-mail your request, including your address to: *homerm@mail.nih.gov.*

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

Dated: December 20, 2010.

Lynn Hellinger,

Director of Management, National Institutes of Health.

[FR Doc. 2010–32911 Filed 12–29–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Mental Health; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Mental Health Initial Review Group; Interventions Committee for Disorders Involving Children and Their Families.

Date: January 31, 2011.

Time: 8:30 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852. (Telephone Conference Call).

Contact Person: David I. Sommers, PhD, Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, National Institutes of Health, 6001 Executive Blvd., Room 6154, MSC 9606, Bethesda, MD 20892–9606. 301–443–7861. *dsommers@mail.nih.gov.*

Name of Committee: National Institute of Mental Health Initial Review Group; Mental Health Services in Non-Specialty Settings.

Date: February 8, 2011.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Melrose Hotel, 2430 Pennsylvania Ave., NW., Washington, DC 20037.

Contact Person: Aileen Schulte, PhD, Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH, Neuroscience Center, 6001 Executive Blvd, Room 6140, MSC 9608, Bethesda, MD 20892–9608. 301–443–1225. *aschulte@mail.nih.gov.*

Name of Committee: National Institute of Mental Health Initial Review Group; Interventions Committee for Adult Disorders.

Date: February 8, 2011.

Time: 8:30 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852. (Telephone Conference Call).

Contact Person: David I. Sommers, PhD, Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, National Institutes of Health, 6001 Executive Blvd., Room 6154, MSC 9606, Bethesda, MD 20892–9606. 301–443–7861. *dsommers@mail.nih.gov.*

Name of Committee: National Institute of Mental Health Initial Review Group; Mental Health Services in MH Specialty Settings.

Date: February 11, 2011.

Time: 8:30 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Melrose Hotel, 2430 Pennsylvania Ave., NW., Washington, DC 20037.

Contact Person: Marina Broitman, PhD, Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH, Neuroscience Center, 6001 Executive Blvd., Room 6153, MSC 9608, Bethesda, MD 20892–9608. 301–402–8152. *mbroitma@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.242, Mental Health Research Grants; 93.281, Scientist Development Award, Scientist Development Award for Clinicians, and Research Scientist Award; 93.282, Mental Health National Research Service Awards for Research Training, National Institutes of Health, HHS)

Dated: December 23, 2010.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–32907 Filed 12–29–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Substance Abuse Prevention; Notice of Meeting

Pursuant to Public Law 92–463, notice is hereby given of the meeting of the Substance Abuse and Mental Health Services Administration (SAMHSA) Center for Substance Abuse Prevention Drug Testing Advisory Board (DTAB) on January 26 and 27, 2011.

A portion of the meeting will be open and will include the Federal drug testing updates from the Department of Transportation, the Department of Defense, the Nuclear Regulatory Commission, and the Federal Drug-Free Workplace Programs; updates on the Mandatory Guidelines for Federal Workplace Drug Testing Programs (the Guidelines); review of the topics that the DTAB will be addressing in the future, including alternate matrices, the electronic custody and control form, and the medical review officer certification; a historical perspective of oral fluid as a drug testing matrix; and the current perspective of the oral fluid matrix, including specimen, drug analytes and their cutoffs, methodologies, proficiency testing, best practices experiences, and specimen drug testing data.

The public is invited to attend the open session in person or to listen via teleconference. Due to the limited space, attendance will be on a registration-only basis. Public comments are welcome. To register, to make arrangements to attend, to obtain the teleconference call-in numbers and access codes, to submit