within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget, Paperwork Reduction Project, *Fax:* 202–395–7285, *E-mail: OIRA_SUBMISSION@OMB.EOP.GOV*, *Attn:* Desk Officer for the Administration for Children and Families.

Robert Sargis.

Reports Clearance Officer.

[FR Doc. 2011–15958 Filed 6–24–11; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0471]

2011 Scientific Meeting of the National Antimicrobial Resistance Monitoring System; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

The Food and Drug Administration (FDA) is announcing a public meeting entitled: ''2011 Scientific Meeting of the National Antimicrobial Resistance Monitoring System.'' The topic to be discussed is animal and retail sampling methods for the National Antimicrobial Resistance Monitoring System (NARMS).

Date and Time: The public meeting will be held on July 20, 2011, from 8 a.m. to 5 p.m.

Location: The public meeting will be held at Holiday Inn Select St. Louis Downtown Convention Center Hotel, 811 North 9th Street, St. Louis, MO 63101, 314–421–4000, FAX: 314–421–5974.

Contact Person: Aleta Sindelar, Center for Veterinary Medicine (HFV–3), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9004, FAX: 240–276–9001, e-mail: *Aleta.Sindelar@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The main purpose of the meeting is to explore ways in which NARMS can improve sampling using current resources. Other topics include:

(1) How should NARMS define adequate sampling for resistance trends?

(2) What are some additional sources for unbiased food animal samples?

(3) What additional information should NARMS collect and report?

Requests for Oral Presentations: Interested persons may present data, information, or views, orally or in writing, on the topic of the discussion of the meeting. Written submissions may be made to the contact person on or before July 6, 2011. Oral presentations from the public during the open public comment period will be scheduled between approximately 2 and 3 p.m. on July 20, 2011. Those desiring to make oral presentations should notify the contact person by July 6, 2011, and submit a brief statement of the general nature of information they wish to present and an indication of the approximate time requested to make their presentation. Time allotted for each presentation may be limited. The contact person will inform each speaker of their schedule prior to the meeting.

Registration is not required for this meeting, however, early arrival is recommended because seating may be limited.

If you need special accommodations due to a disability, please contact Aleta Sindelar (see *Contact Person*) at least 7 days in advance.

Comments: Regardless of attendance at the public meeting, interested persons may submit to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, either electronic or written comments regarding this document. Submit electronic comments to *http://www.regulations.gov.* It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. The docket will remain open for written or electronic comments for 30 days following the meeting.

Agenda: The meeting will address goals and challenges of surveying retail meats and food animals for antimicrobial susceptibility in foodborne bacteria. The agenda for the public meeting will be made available on the Agency's Web site at *http://www.fda.gov/AnimalVeterinary/SafetyHealth/AntimicrobialResistance/NationalAntimicrobialResistanceMonitoringSystem/ucm059135.htm.*

Transcripts: FDA will prepare a meeting transcript and make it available on the Agency's Web site (*see Agenda*) after the meeting. FDA anticipates that transcripts will be available approximately 60 business days after the meeting. The transcript will be available for public examination at the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday. A transcript will also be available in either hardcopy or on CD–ROM after submission of a Freedom of Information request. Written requests are to be sent to Division of Freedom of Information (ELEM–1029), Food and Drug Administration, 12420 Parklawn Dr., Element Bldg. Rockville, MD 20857.

Dated: June 16, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–15982 Filed 6–24–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB review; comment request Health Information National Trends Survey 4 (HINTS 4) (NCI)

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute (NCI), the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on April 22, 2011 (76 FR 22714) and allowed 60-days for public comment. One public comment was received on April 23, 2011 which commented on the number of previous surveys and expense. An e-mail response was sent on April 25, 2011, stating, ''Thank you for your comments. We will take your comments into consideration.'' The purpose of this notice is to allow an additional 30 days for public comment. 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: Health Information National Trends Survey 4 (HINTS 4) (NCI) (OMB 0925–0538, Exp 11/30/2008). *Type of Information Collection Request: Reinstatement with Change. Need and Use of Information Collection:* HINTS 4 will provide NCI with a comprehensive assessment of the American public's current access to, and

use of, information about cancer across the cancer care continuum from cancer prevention, early detection, diagnosis, treatment, and survivorship. The content of the survey will focus on understanding the degree to which members of the general population understand vital cancer prevention messages. More importantly, this NCI survey will couple knowledge-related questions with inquiries into the communication channels through which understanding is being obtained, and assessment of cancer-related behavior. The Public Health Services Act, Sections 411 (42 U.S.C. 285a) and 412 (42 U.S.C. 285a-1.1 and 285a-1.3), outline the research and information dissemination mission of the NCI which authorizes the collection of this information. *Frequency of Response:* Once. *Affected Public:* Individuals. *Type of Respondents:* U.S. adults (persons aged 18+). The annual reporting burden is documented in the table below. There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

Data collection cycle	Type of respondent	Number of respondents	Frequency of response	Average time per response minutes/ hour	Annual hour burden
Cycle 1	Mail survey	3,533	1	30/60 (.5)	1,766.5
Cycle 2	Mail survey	3,533	1	30/60 (.5)	1,766.5
Cycle 3	Mail survey	3,500	1	30/60 (.5)	1,750
Cycle 4	Mail survey	3,500	1	30/60 (.5)	1,750
Total					7,033

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:* NIH Desk Officer, Office of Management and Budget, at *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Bradford W. Hesse, PhD, Project Officer, National Cancer Institute, NIH, EPN 4068, 6130 Executive Boulevard, MSC 7365, Bethesda, Maryland 20892–7365, or call non-toll free number 301–594–9904 or fax your request to 301–480–2198, or e-mail your request, including your address, to *hesseb@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: June 20, 2011.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2011–15994 Filed 6–24–11; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2) notice is hereby given of the following meeting.

The meeting 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 purpose of this meeting is to evaluate requests for preclinical development resources, biologics, clinical assays and other developmental programs for potential new therapeutics for the treatment of cancer. The outcome of the evaluation will provide information to internal NCI committees that will decide whether NCI should support requests and make available contract resources for development of the potential therapeutic to improve the treatment of various forms of cancer. The research proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material and personal information concerning individuals associated with the proposed research projects, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Cancer Institute Special Emphasis Panel, Clinical Assay Development Program (CADP).

Date: July 27, 2011.

Time: 9 a.m.–4 p.m.

Agenda: To review grant applications for the CADP.

Place: Bethesda Marriott North Hotel, 5701 Marinelli Road, Bethesda, MD 20852.

Contact Person: Dr. Barbara Conley, Executive Secretary, Clinical Assay Development Program (CADP), National Cancer Institute, NIH, 6130 Executive Boulevard, Room 6035A, Bethesda, MD 20892, 301–496–8639, *conleyba@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)