Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site and call the appropriate advisory committee hotline/phone line to learn about possible modifications before coming to the meeting.

Agenda: On May 11, 2011, the Pediatric Ethics Subcommittee of the Pediatric Advisory Committee will meet to discuss the general topic of the ethics of administering subtherapeutic doses of investigational products to children for the purpose of determining, for example, drug metabolism, disposition, and targeting (*e.g.,* exploratory investigational new drug (IND) studies). In this context, the subcommittee will also discuss the referral of such protocols by an Institutional Review Board for review by a Federal panel under 21 CFR 50.54.

The subcommittee's recommendations will then be presented to the FDA Pediatric Advisory Committee on Monday, May 16, 2011. The announcement of the May 16, 2011, Pediatric Advisory Committee meeting can be found elsewhere in this issue of the **Federal Register**.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm*. Scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the subcommittee. Written submissions may be made to the contact person on or before April 28, 2011. Oral presentations from the public will be scheduled between approximately 11 a.m. and 12 noon on May 11, 2011. Those desiring to make formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before April 20, 2011. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by April 21, 2011.

Comments: FDA is opening a docket to allow for additional public comments to be submitted to the Agency on issues before the Pediatric Ethics Subcommittee beginning April 15, 2011, and closing May 5, 2011. All comments received on or before May 5, 2011, will be provided to the committee members. All comments received after May 5, 2011, will be taken into consideration by the Agency. Interested persons are encouraged to use the docket to submit either electronic or written comments regarding this meeting (*see* **ADDRESSES**). Submit electronic comments to *http://www.regulations.gov*. Submit written comments to Division of Dockets Management (*see* **ADDRESSES**). It is necessary to submit only 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.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Walter Ellenberg at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: April 11, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–9149 Filed 4–14–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Food Reporting Comparison Study (FORCS) and Food and Eating Assessment Study (FEAST) (NCI)

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: Food Reporting Comparison Study (FORCS) and Food and Eating Assessment Study (FEAST) (NCI*). Type of Information Collection Request:* Extension. *Need and Use of Information Collection:* The title of this collection was previously, "24-hour Dietary Recall Method Comparison and the National Cancer Institute (NCI) Observational Feeding Studies." The objective of the two studies is to compare the performance of the newly developed computerized Automated Self-Administered 24-Hour Recall (ASA24) approach to collecting 24 hour recall (24HR) data with the current standard, the interviewer-administered Automated Multiple Pass Method (AMPM). The ultimate goal is to determine to what extent the new automated instrument can be used instead of the more expensive interviewer-administered instrument in the collection of dietary intake data. *Frequency of Response:* Twice. *Affected Public:* Individuals. *Type of Respondents:* For the FORCS study, approximately 1,200 adult members from three health maintenance organization plans (in Minnesota, California, and Michigan) between ages 20 and 70 years. For the FEAST study, approximately 90 adult residents from the Washington, DC metropolitan area between ages 20 and 70 years. The annual reporting burden is estimated at 866 hours (see table below). This amounts to an estimated 2,598 burden hours over the 3-year data collection period with a total cost to the respondents of $54,293. There are no Capital costs, Operating costs, and/or Maintenance Costs to report.

Participants and study	Questionnaire	Number of respondents	Frequency of response	Average time per response minutes/hour	Annual hour burden
General Public for FORCS ...	Refusal Reasons and Demographics (Attach 4A, Screen 8).	1,770	1	5/60 (0.083)	148
	Contact Information (Attach 4A, Screen 5).	400	1	5/60 (0.083)	33
	Screener (Attach 5)	400	1.00	5/60 (0.083)	33
	AMPM (Attach 1)	400	1.00	30/60 (0.50)	200
	ASA24 (Attach 2)	400	1.00	30/60 (0.50)	200
	Demographics and Health Questionnaire (Attach 6).	360	1.00	10/60 (0.167)	60
	Demographics, Health and Preference Questionnaire (Attach 7).	360	1.00	15/60 (0.25)	90
General Public for FEAST	Screener (Attach 8)	33	1.00	5/60 (0.083)	6
	Reminder Telephone Call (Attach 10).	33	1.00	5/60 (0.083)	6
	Eating 3 meals	33	1.00	135/60 (2.25)	151
	Either AMPM or ASA24 (Attach 1 or 2).	33	1.00	30/60 (0.50)	34
	Demographics and Health Questionnaire (Attach 12).	33	1.00	10/60 (0.167)	11
	.....	3,485	.....	.....	866

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) 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) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to 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.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans, contact Frances E. Thompson, PhD, Project Officer, National Cancer Institute, NIH, EPN 4095A, 6130 Executive Boulevard MSC 7335, Bethesda, Maryland 20892–7335, or call non-toll-free number 301–594–4410, or FAX your request to 301–435–3710, or e-mail your request, including your address, to *thompsof@mail.nih.gov.*

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60-days of this publication.

Dated: April 8, 2011.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2011–9204 Filed 4–14–11; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meeting

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

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the 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 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: Council of Councils.

Date: June 29, 2011.

Open: 8:30 a.m. to 12:30 p.m.

Agenda: Call to Order and Introductions; Announcements and Future Meeting Dates; DPCPSI Update; Remarks by the Director, NIH; and Review of NIH Common Fund Initiative Concepts: Process and Criteria.

Place: National Institutes of Health, Building 31, 31 Center Drive, Conference Room 6, Bethesda, MD 20892.

Closed: 1:30 p.m. to 2:20 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Building 31, 31 Center Drive, Conference Room 6, Bethesda, MD 20892.

Open: 2:20 p.m. to 5:30 p.m.

Agenda: Overall Discussion & Recommendations and Closing Remarks.

Place: National Institutes of Health, Building 31, 31 Center Drive, Conference Room 6, Bethesda, MD 20892.

Contact Person: Robin Kawazoe, Executive Secretary, Division Of Program Coordination, Planning, And Strategic Initiatives, Office Of The Director, NIH, Building 1, Room 260B, Bethesda, MD 20892. *kawazoer@mail.nih.gov.*

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person. Information is also available on the Council of Council's home page at *http://dpcpsi.nih.gov/council/,* where an agenda and proposals to be discussed will be posted before the meeting date.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license,