One collection, entitled ''Health and Diet Survey—General Topics,'' tracks a broad range of consumer attitudes, awareness, knowledge, and self-reported behaviors related to key diet and health issues. The other collection, entitled ''Health and Diet Survey—*Dietary Guidelines* Supplement,'' will provide FDA with updated information about consumer attitudes, awareness, knowledge, and behavior regarding various elements of nutrition and physical activity based on the key recommendations of the *Dietary Guidelines for Americans*, which are jointly issued by the Department of Health and Human Services and Department of Agriculture every 5 years.

The information to be collected with the Health and Diet Survey—General Topics will include: (1) Awareness of diet-disease relationships; (2) food and dietary supplement label use; (3) dietary practices including strategies to lose or maintain weight; and (4) awareness and knowledge of dietary fats. The information to be collected with the Health and Diet Survey—*Dietary Guidelines* Supplement will include: (1) Opinions about the nutrition information provided by the government; (2) awareness and familiarity with government nutrition programs and publications such as the Food Guide Pyramid and the *Dietary Guidelines for Americans*; (3) knowledge of the relationships between food choices, exercise habits, weight loss, and health; (4) choices surrounding exercise, calorie intake, saturated and trans fats, fruits and vegetables, whole grains, dairy, fish, meat, cholesterol, carbohydrates, salt, and sugar. The survey will also ask about use of Federal nutrition information, special diet, weight status, health status, and demographics.

FDA and other Federal agencies will use the information from the Health and Diet Survey to evaluate and develop strategies and programs to encourage and help consumers adopt healthy lifestyles. The information will also help FDA and other Federal agencies evaluate and track consumer awareness and behavior as outcome measures of their achievement in improving public health.

Description of Respondents: The respondents are adults, age 18 and older, drawn from the 50 states and the District of Columbia. Participation will be voluntary.

In the **Federal Register** of May 25, 2007 (72 FR 29332), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Activity	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
General Topics: Pretest	27	1	27	0.25	6.75
General Topics: Screener	10,000	1	10,000	0.02	200
General Topics: Survey	3,000	1	3,000	0.25	750
Dietary Guidelines Supplement: Screener	4,000	1	4,000	0.02	80
Dietary Guidelines Supplement: Survey	1,200	1	1,200	0.22	264
Total					1,300.75

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA has based its estimate of the number of respondents and the burden hours per response on its experience with the Health and Diet Survey over the past 3 years. The agency will use a screener to select an eligible adult respondent in each household to participate in the survey. For the Health and Diet Survey—General Topics data collection activity a total of 3,000 adults in the 50 states and the District of Columbia will be interviewed by telephone. We estimate that it will take a respondent 1.2 minutes (0.02 hours) to complete the screening questions and 15 minutes (0.25 hours) to complete the entire survey. Prior to the administration of the survey, the agency plans to conduct a pretest to identify and resolve potential problems. The pretest will be conducted with 27 participants; we estimate that it will take a respondent 15 minutes (0.25 hours) to complete the pretest. For the Health and Diet Survey—*Dietary Guidelines*Supplement data collection activity a total of 1,200 adults in the 50 states and the District of Columbia will be interviewed by telephone. We estimate that it will take a respondent 1.2 minutes (0.02 hours) to complete the screening questions and 13.2 minutes (0.22 hours) to complete the entire survey. Target sample size of the combined data collection is 4,200 respondents who complete the survey.

Dated: December 7, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–24123 Filed 12–12–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Quality System Regulation Educational Forum on Design Controls; Public Workshop; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an amendment to the notice of public workshop entitled ''Quality System Regulation Educational Forum on Design Controls.'' This workshop was announced in the **Federal Register** of October 11, 2007 (72 FR 57951). The amendment is made to reflect a change in the *Location* portion of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT: David Arvelo, Food and Drug

Administration, 4040 North Central Expressway, suite 900, Dallas, TX 75204, 214–253–4952, FAX: 214–253–4970, e-mail: *david.arvelo@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of October 11, 2007 (72 FR 57951), FDA announced that a public workshop entitled ''Quality System Regulation Educational Forum on Design Controls'' would be held on Friday, April 4, 2008. On page 57951, in the second column, the *Location* portion of the document is amended to read as follows:

Location: The public workshop will be held at the Adam's Mark Hotel Dallas, 400 North Olive St., Dallas, TX 75201, 214–922–8000. Directions to the facility and additional information are available at the FDA Medical Device Industry Coalition Web site at *http://www.fmdic.org/*.

Dated: December 7, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–24144 Filed 12–12–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of Closed Meetings

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 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/or contract proposals 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 and/or contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Cancer Institute Special Emphasis Panel, Operations and Technical Support at the NCI-Frederick.

Date: January 7, 2008.

Time: 8 a.m. to 4 p.m.

Agenda: To review and evaluate contract proposals.

Place: Gaithersburg Hilton, 620 Perry Parkway, Gaithersburg, MD 20877.

Contact Person: Lalita D. Palekar, PhD. Scientific Review Administrator, Special Review and Logistics Branch, Division of Extramural Activities, National Cancer Institute, 6116 Executive Blvd., Bethesda, MD 20892–7405, 301–496–7575, *palekarl@mail.nih.gov.*

Name of Committee: National Cancer Institute Initial Review Group, Subcommittee G–Education.

Date: February 12–13, 2008.

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

Agenda: To review and evaluate grant applications.

Place: Crowne Plaza Washington Silver Spring, 8777 Georgia Ave., Silver Spring, MD 20910.

Contact Person: Sonya Roberson, PhD, Scientific Review Administrator, Resources and Training Review Branch, Division of Extramural Activities, National Cancer Institute, 6116 Executive Blvd., Room 8109, Bethesda, MD 20892, 301–594–1182, *robersos@mail.nih.gov.*

Name of Committee: National Cancer Institute Special Emphasis Panel, R25(E) Special Emphasis Panel (SEP).

Date: February 12, 2008.

Time: 5 p.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: Crowne Plaza Washington Silver Spring, 8777 Georgia Ave., Silver Spring, MD 20910.

Contact Person: Lynn M. Amende, PhD, Scientific Review Administrator, Resources and Training Review Branch, Division of Extramural Activities, National Cancer Institute, 6116 Executive Boulevard Room 8105 Bethesda, MD 20892–8328, 301–451–4759, *amendel@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)

Dated: December 4, 2007.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 07–6034 Filed 12–12–07; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Biomedical Imaging and Bioengineering; Notice of 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 a meeting of the National Advisory Council for Biomedical Imaging and Bioengineering.

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/or contract proposals 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 and/or contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Advisory Council for Biomedical Imaging and Bioengineering, NACBIB January 2008.

Date: January 25, 2008.

Open: 8:30 a.m. to 1 p.m.

Agenda: Report from the Institute Director and other Institute staff and presentations of working group reports.

Place: Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817.

Closed: 1 p.m. to 4:30 p.m.

Agenda: To review and evaluate grant applications and/or proposals.

Place: Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817.

Contact Person: Anthony Demsey, PhD, Director, National Institute of Biomedical Imaging and Bioengineering, 6701 Democracy Blvd., Room 241, Bethesda, MD 20892.

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 Institute's/Center's home page: *http://www.nibib1.nih.gov/about/NACBIB/NACBIB.htm*, where an agenda and any additional information for the meeting will be posted when available.

Dated: December 5, 2007.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 07–6031 Filed 12–12–07; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute On Aging; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as