

notification of attendance is June 10, 2011.

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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: May 16, 2011.

**Elaine L. Baker,**

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

[FR Doc. 2011-12562 Filed 5-20-11; 8:45 am]

**BILLING CODE 4163-18-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

The meeting announced below concerns Centers for Agricultural Disease and Injury Research, Education, and Prevention, Program Announcement (PA) Number PAR-11-022, initial review.

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

#### TIME AND DATE:

8 a.m.-6:30 p.m., June 6, 2011  
(Closed)

8 a.m.-6:30 p.m., June 7, 2011  
(Closed)

8 a.m.-6:30 p.m., June 8, 2011  
(Closed)

8 a.m.-6:30 p.m., June 9, 2011  
(Closed)

8 a.m.-6:30 p.m., June 10, 2011  
(Closed)

*Place:* Doubletree by Hilton Philadelphia Center City, 237 S. Broad Street, Philadelphia, Pennsylvania 19107.

*Status:* The meeting 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 Public Law 92-463.

*Matters To Be Discussed:* The meeting will include the initial review, discussion, and evaluation of applications received in response to "Centers for Agricultural Disease and

Injury Research, Education, and Prevention, PAR-11-022, initial review."

*Contact Person for More Information:* M. Chris Langub, PhD, Scientific Review Officer, Office of Extramural Programs, 1600 Clifton Road, NE., Mailstop E74, Atlanta, Georgia 30333, Telephone: (404) 498-2543.

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Dated: May 16, 2011.

**Elaine L. Baker,**

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

[FR Doc. 2011-12561 Filed 5-20-11; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Healthcare Infection Control Practices Advisory Committee (HICPAC)

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

#### TIMES AND DATES:

9 a.m.-5 p.m., June 16, 2011.

9 a.m.-12 p.m., June 17, 2011.

*PLACE:* CDC, Global Communications Center, Building 19, Auditorium B3, 1600 Clifton Road, NE., Atlanta, Georgia, 30333.

*STATUS:* Open to the public, limited only by the space available. Please register for the meeting at <http://www.cdc.gov/hicpac>.

*PURPOSE:* The Committee is charged with providing advice and guidance to the Secretary, the Assistant Secretary for Health, the Director, CDC, the Director, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), and the Director, Division of Healthcare Quality Promotion regarding (1) The practice of healthcare infection control; (2) strategies for surveillance, prevention, and control of infections (e.g., nosocomial infections), antimicrobial resistance, and related events in settings where healthcare is provided; and (3) periodic updating of

guidelines and other policy statements regarding prevention of healthcare-associated infections and healthcare-related conditions.

**MATTERS TO BE DISCUSSED:** The agenda will include updates on CDC's activities for healthcare associated infections; draft guidelines for prevention of infections among patients in neonatal intensive care units (NICU); draft guidelines for infection control in healthcare personnel; draft guidelines for the prevention of surgical site infections; update from the HICPAC surveillance working group; and discussion of a draft infection control worksheet for acute-care hospitals. Agenda items are subject to change as priorities dictate.

#### CONTACT PERSON FOR MORE INFORMATION:

Heidi Williams, HICPAC, Division of Healthcare Quality Promotion, NCEZID, CDC, 1600 Clifton Road, NE., Mailstop A-07, Atlanta, Georgia 30333, Telephone (404) 639-4227. Email: [hicpac@cdc.gov](mailto:hicpac@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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: May 16, 2011.

**Elaine L. Baker,**

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

[FR Doc. 2011-12560 Filed 5-20-11; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

#### Office of Planning, Research and Evaluation; Advisory Committee on Head Start Research and Evaluation

**AGENCY:** Administration for Children and Families, Department of Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** This notice announces a forthcoming meeting of a public advisory committee of the Administration for Children and Families (ACF). The meeting will be open to the public.

*Name of Committee:* Advisory Committee for Head Start Research and Evaluation.

*General Function of Committee:* The Advisory Committee for Head Start Research and Evaluation will provide feedback on the published final report for the Head Start Impact Study, offering interpretations of the findings, discussing implications for practice and policy, and providing recommendations on follow-up research, including additional analysis of the Head Start Impact Study data. The Committee will also be asked to provide recommendations to the Secretary regarding how to improve Head Start and other early childhood programs by enhancing the use of research-informed practices in early childhood. Finally, the Committee will be asked to provide recommendations on the overall Head Start research agenda, including—but not limited to—how the Head Start Impact Study fits within this agenda. The Committee will provide advice regarding future research efforts to inform HHS about how to guide the development and implementation of best practices in Head Start and other early childhood programs around the country.

**DATES:** The meeting will be held from 8:30 a.m. to 5 p.m. on June 7–8, 2011.

**ADDRESSES:** The Madison Hotel, 1177 15th Street, NW., Washington, DC 20005. Phone: (202) 862-1600.

**FOR FURTHER INFORMATION CONTACT:** Jennifer Brooks, Office of Planning, Research, and Evaluation, e-mail [jennifer.brooks@acf.hhs.gov](mailto:jennifer.brooks@acf.hhs.gov) or call (202) 205-8212.

*Agenda:* The Committee will review information on the Federal and Early Head Start programs and the children and families they serve, and learn about the latest research in the area of parent, family, and community engagement and other topic areas related to early childhood education and development.

*Procedure:* Interested persons may present data, information or views, in writing, on issues pending before the Committee. Written submissions may be made to the contact person on or before May 24, 2011. All written materials provided to the contact person will be shared with the Committee members.

ACF welcomes the attendance of the public at this advisory committee meeting 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 Jennifer Brooks at least seven days in advance of the meeting. Information about the Committee and this meeting can be found at the Committee Web site, [http://www.acf.hhs.gov/prosrams/opre/hs/advisory\\_com/](http://www.acf.hhs.gov/prosrams/opre/hs/advisory_com/).

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

Dated: May 4, 2011.

**David A. Hansell,**  
*Acting Assistant Secretary for Children and Families.*

[FR Doc. 2011-12370 Filed 5-20-11; 8:45 am]

**BILLING CODE M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2011-N-0001]

#### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Health and Diet Survey

**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 June 22, 2011.

**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: 202-395-7285, or e-mailed to [oir\\_submission@omb.eop.gov](mailto:oir_submission@omb.eop.gov). All comments should be identified with the OMB control number 0910-0545. Also include the FDA docket number found in brackets in the heading of this document.

**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 compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

#### Health and Diet Survey—(OMB Control Number 0910-0545)—Extension

FDA is seeking extension of OMB approval for the Health and Diet Survey, which is a voluntary consumer survey

intended to gauge and track consumer attitudes, awareness, knowledge, and behavior regarding various topics related to health, nutrition, and physical activity. The authority for FDA to collect the information derives from FDA's Commissioner of Food and Drugs authority provided in section 903(d)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(d)(2)).

The survey consists of two independent data collection activities. 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 the U.S. 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. This survey has been repeated approximately every 3 years over the course of the past several years for the purpose of tracking changes and trends in public opinions and consumer behavior, with some new questions added or omitted or partially modified each iteration in response to current events. In the next 3 years, FDA plans to field the Health and Diet Survey—General Topics in 2012 and anticipates that it might have the need for additional iterations in 2014. The information to be collected with the Health and Diet Survey—*Dietary Guidelines* Supplement will include: (1) Awareness and sources of information, (2) attitudes toward diet and physical activity, and (3) practice and knowledge related to recommended behaviors. The survey will also ask about perceptions and use of Federal nutrition information, special diet, weight status, health status, and demographics. In the next 3 years, FDA anticipates to field the Health and Diet Survey—*Dietary Guidelines* Supplement in 2011–2012.

FDA and other Federal Agencies will use the information from the Health and Diet Survey to evaluate and develop strategies and programs to encourage