

(Greentree), Pittsburgh, PA 15220. This meeting will also be available by remote access. Registration information is available on the NIOSH Web site at <http://www.cdc.gov/niosh>.

**FOR FURTHER INFORMATION CONTACT:**

David Book, NIOSH National Personal Protective Technology Laboratory (NPPTL), 626 Cochran Mill Road, Pittsburgh, PA 15236 (412) 386-6691 or (412) 386-5200 (these are not toll free numbers).

**SUPPLEMENTARY INFORMATION:**

**Public Meeting**

NIOSH will hold a public meeting to allow commenters to present information on an increase in respirator certification and approval fees on individual respirator manufacturers, the respirator market, or on those industries that rely on NIOSH approved respiratory equipment.

Requests to make presentations at the public meeting should be mailed to the NIOSH Docket Office, Robert A. Taft Laboratories, MS-C34, 4676 Columbia Parkway, Cincinnati, OH 45226. Requests may also be submitted by telephone (513) 533-8611, facsimile (513) 533-8285, or emailed to [nioshdocket@cdc.gov](mailto:nioshdocket@cdc.gov) with the words "respirator fees presentation" in the subject line. All requests to present should contain the name, address, telephone number and relevant business affiliations of the presenter, and the approximate time requested for the presentation. Oral presentations will be limited to 15 minutes. After reviewing the requests for presentations, NIOSH will notify the presenter that his/her presentation is scheduled. If a participant is not in attendance when his/her presentation is scheduled to begin, the remaining participants will be heard in order. After the last scheduled speaker is heard, participants who missed their assigned times may be allowed to speak, limited by time available.

Attendees who wish to speak but did not submit a request for the opportunity to make a presentation may be given this opportunity after the scheduled speakers are heard, at the discretion of the presiding officer and limited by time available. This meeting will also be using audio/LiveMeeting Conferencing, remote access capabilities where interested parties may listen in and view the presentations over the Internet simultaneously. Parties remotely accessing the meeting will have the opportunity to comment during the open comment period.

Registration is required for both in-person and LiveMeeting participation.

Because this meeting is being held at a Federal site, pre-registration is required on or before April 26, 2013 and a government-issued photo ID (driver's license or passport) will be required to obtain entrance to the facility. Non-US citizens need to register by March 29, to allow sufficient time for mandatory facility security clearance procedures to be completed.

An email confirming registration will be sent from NIOSH for both in-person participation and audio conferencing participation. Details required to participate via the conferencing will be provided by NIOSH in a separate email. This option will be available to participants on a first come, first served basis and is limited to the first 100 participants.

Registration information is available on the NIOSH Web site at <http://www.cdc.gov/niosh>.

Dated: March 20, 2013.

**John Howard,**

*Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.*

[FR Doc. 2013-06850 Filed 3-26-13; 8:45 am]

**BILLING CODE 4163-19-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)**

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 of the aforementioned committee.

*Time and Date:* 8:30 a.m.–3:00 p.m. (EDT), April 25, 2013.

*Place:* CDC, Building 21, Rooms 1204 A/B, 1600 Clifton Road, NE., Atlanta, Georgia 30333.

*Status:* Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people. The public is welcome to participate during the public comment period, which is tentatively scheduled from 2:45 p.m. to 2:50 p.m. This meeting is also available by teleconference. Please dial (877) 930-8819 and enter code 1579739.

*Purpose:* The committee will provide advice to the CDC Director on strategic and other broad issues facing CDC.

*Matters To Be Discussed:* The Advisory Committee to the Director will

receive updates from the Global Workgroup; State, Tribal, Local and Territorial Workgroup; the Communications Workgroup; the Ethics Subcommittee, and the Health Disparities Subcommittee, as well as an update from the CDC Director.

Agenda items are subject to change as priorities dictate.

*Contact Person for More Information:* Carmen Villar, M.S.W., Designated Federal Officer, Advisory Committee to the Director, CDC, 1600 Clifton Road NE., M/S D-14, Atlanta, Georgia 30333. Telephone 404/639-7000, Email: [GHickman@cdc.gov](mailto:GHickman@cdc.gov). The deadline for notification of attendance is April 22, 2013. To register for this meeting, please send an email to [ACDDirector@cdc.gov](mailto:ACDDirector@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.

**Dana Redford,**

*Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

[CFDA Number: 93.676]

**Announcement of the Award of Fifteen Single-Source Program Expansion Supplement Grants to Unaccompanied Alien Children's Shelter Care Grantees**

**AGENCY:** Office of Refugee Resettlement, Administration for Children and Families, Department of Health and Human Services.

**ACTION:** Announcement of the award of fifteen single-source program expansion grants to ten current grantees to expand bed capacity and supportive services to the increasing number of unaccompanied alien children.

**SUMMARY:** The Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR) announces the award of fifteen single-source program expansion supplement grants to the following ten current grantees, for a total of \$47,168,490.

Organization	Location	Amount
BCFS Health and Human Services .....	San Antonio, TX .....	\$3,039,665
Heartland Human Care Services, Inc. ....	Chicago, IL .....	1,659,393
Southwest Key, Inc. ....	Austin, TX .....	13,431,660
Children's Village .....	Dobbs Ferry, NY .....	4,208,741
United States Conference of Catholic Bishops .....	Washington, DC .....	209,576
Lutheran Immigration Refugee Services .....	Baltimore, MD .....	281,452
US Committee for Refugees and Immigrants .....	Washington, DC .....	602,690
International Education Services .....	Los Fresnos, TX .....	16,314,360
Lincoln Hall .....	Lincolndale, NY .....	7,024,414
David and Margaret Youth & Family Services .....	LaVerne, CA .....	396,539

These supplement grants will support the expansion of bed capacity and supportive services to meet the number of unaccompanied alien children referrals from the Department of Homeland Security (DHS). The funding program is mandated by section 462 of the Homeland Security Act to ensure appropriate placement of all referrals from the DHS. The program is tied to DHS apprehension strategies and sporadic number of border crossers. Award funds will support services to unaccompanied alien children through September 30, 2013.

**DATES:** The period of support under these supplements is October 1, 2012 through September 30, 2013.

**FOR FURTHER INFORMATION CONTACT:** Jallyn Sualog, Acting Director, Division of Children's Services, Office of Refugee Resettlement, 901 D Street SW., Washington, DC 20447, Telephone (202) 401-4997. Email: [jallyn.sualog@acf.hhs.gov](mailto:jallyn.sualog@acf.hhs.gov)

**SUPPLEMENTARY INFORMATION:** Since the beginning of FY 13, the Unaccompanied Alien Children (UAC) program has seen a dramatic increase in the number of DHS referrals. The influx of border crossers referred by DHS has grown beyond anticipated rates and has resulted in the program needing a significant increase in the number of shelter beds and supportive services.

The UAC program has specific requirements for the provision of services to unaccompanied alien children. These grantee organizations are the only entities with the infrastructure, licensing, experience, and appropriate level of trained staff to meet the required service requirements and the urgent need for the expansion of services required to respond to unexpected arrivals of unaccompanied children. The program expansion supplement will support such services and alleviate the buildup of children waiting in border patrol stations for placement in shelter care.

**Statutory Authority:** Section 462 of the Homeland Security Act, (6 U.S.C. 279) and sections 235(c) and 235(d) of the William

Wilberforce Trafficking Victims Protection Reauthorization Act of 2008, (8 U.S.C. 1232(c) and 1232(d)).

**Eskinder Negash,**  
*Director, Office of Refugee Resettlement.*  
[FR Doc. 2013-07061 Filed 3-26-13; 8:45 am]  
**BILLING CODE 4184-45-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2013-N-0065]

#### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Registration of Food Facilities 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 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 April 26, 2013.

**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 emailed to [oir\\_submission@omb.eop.gov](mailto:oir_submission@omb.eop.gov). All comments should be identified with the OMB control number 0910-0502. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Domini Bean, Office of Information Management, Food and Drug

Administration, 1350 Piccard Dr., PI50-400T, Rockville, MD 20850, 301-796-5733, [domini.bean@fda.hhs.gov](mailto:domini.bean@fda.hhs.gov).

**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.

#### Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002—21 CFR 1.230-1.235 (OMB Control Number 0910-0502)—Extension

The Public Health Security and Bioterrorism Preparedness and Response Act of 2002 (the Bioterrorism Act) (Pub. L. 107-188) added section 415 to the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 350d), which requires domestic and foreign facilities that manufacture, process, pack, or hold food for human or animal consumption in the United States to register with the Food and Drug Administration (FDA). Sections 1.230 through 1.235 of FDA's regulations (21 CFR 1.230-1.235) set forth the procedures for registration of food facilities. Information provided to FDA under these regulations helps the Agency to notify quickly the facilities that might be affected by a deliberate or accidental contamination of the food supply. In addition, data collected through registration is used to support FDA enforcement activities and to screen imported food shipments. Advance notice of imported food allows FDA, with the support of the Bureau of Customs and Border Protection, to target import inspections more effectively and help protect the nation's food supply against terrorist acts and other public health emergencies. If a facility is not registered or the registration for a facility is not updated when necessary, FDA may not be able to contact the facility and may not be able to target import inspections effectively in case of a known or potential threat to the food supply or other food-related emergency, putting consumers at risk of consuming hazardous food products that could