participating in three focus groups (in the form of safety circle workgroup meetings) that will take approximately 60 minutes. The 60 minutes includes a 30 minute discussion and the completion of a focus group worksheet and at one point, a dust control worksheet. The focus groups will debrief general CPDM data so participants can dialogue about ways to lower their exposure levels. In addition, workers will be asked to complete a pre and post survey (~15 minutes). It also is estimated that a sample of up to nine mine site leaders will participate in the form of interviews/focus groups about HSMS practices at the same mining operations which have agreed to participate, and complete a dust assessment form. The interviews/focus groups also will occur two to three times during each of the NIOSH field visits and will take no more than 45 minutes each. All participants will be between the ages of 18 and 75, currently employed, and living in the United States. Participation will require no more than 3 hours of workers' time over the six-week intervention and no more than 2.5 hours of site leaders' time over the six-week intervention period.

There is no cost to respondents other than their time. The total burden in time is an estimated 64 burden hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hours)
Mine & Health Safety Managers/Leaders.	Mine Recruitment and Participation Script	3	1	5/60
	Worksite Leadership Interview/Focus Group Guide	3	3	45/60
	Controls to Reduce Respirable Dust Exposure Assessment Worksheet for Workers and Management.	3	1	15/60
Individual Mine Workers	Mine Worker Recruitment Script	50	1	5/60
	Pre/Post Mine Worker Survey	50	2	15/60
	Mine Worker CPDM Focus Group Guide	50	3	30/60
	Controls to Reduce Respirable Dust Exposure Assessment Worksheet for Workers and Management.	50	1	15/60
	Mine Worker Focus Group Worksheet	50	3	15/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2015–13383 Filed 6–1–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Subcommittee for Dose Reconstruction Reviews (SDRR), Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board), National Institute for Occupational Safety and Health (NIOSH)

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 subcommittee:

TIME AND DATE: 10:30 a.m.–5:00 p.m. EDT, June 24, 2015

PLACE: Audio Conference Call via FTS Conferencing.

STATUS: Open to the public, but without a public comment period. The public is welcome to submit written comments in advance of the meeting, to the contact person below. Written comments received in advance of the meeting will be included in the official record of the meeting. The public is also welcome to listen to the meeting by joining the teleconference at the USA toll-free, dial-in number at 1–866–659–0537 and the pass code is 9933701.

BACKGROUND: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines that have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction, which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC).

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2015.

PURPOSE: The Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under E.O. 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee for Dose Reconstruction Reviews was established to aid the Advisory Board in carrying out its duty to advise the Secretary, HHS, on dose reconstruction.

MATTERS FOR DISCUSSION: The agenda for the Subcommittee meeting includes the following dose reconstruction program quality management and assurance activities: Current findings from NIOSH and Advisory Board dose reconstruction blind reviews; dose reconstruction cases under review from Sets 14–18, including the Oak Ridge sites (Y–12, K–25, Oak Ridge National Laboratory, and Savannah River Site; preparation of the Advisory Board's next report to the

Secretary, HHS, summarizing the results of completed dose reconstruction reviews.

The agenda is subject to change as priorities dictate.

CONTACT PERSON FOR MORE INFORMATION: Theodore Katz, Designated Federal Officer, NIOSH, CDC, 1600 Clifton Road, Mailstop E–20, Atlanta GA 30333, Telephone (513)533–6800, Toll Free 1(800) CDC–INFO, Email *ocas@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.

Elaine L. Baker,

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

[FR Doc. 2015–13311 Filed 6–1–15; 8:45 am]

BILLING CODE 4163–19–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

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 a meeting for the initial review of applications in response to Funding Opportunity Announcement (FOA), IP–11–00100501PPHF15, PPHF–2015—Enhanced Surveillance for New Vaccine Preventable Disease—A Program Expansion for Acute Respiratory Illness Surveillance Financed Solely by 2015 Prevention and Public Health Funds.

Time and Date: 10:00 a.m.–3:00 p.m., June 23, 2015 (Closed).

Place: Teleconference.

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 for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to ''PPHF–2015-Enhanced Surveillance for New Vaccine Preventable Disease—A Program Expansion for Acute Respiratory Illness Surveillance Financed Solely by 2015 Prevention and Public Health Funds'', IP–11–00100501PPHF15.

Contact Person for More Information: Gregory Anderson, M.S., M.P.H., Scientific Review Officer, CDC, 1600 Clifton Road NE., Mailstop E60, Atlanta, Georgia 30333, Telephone: (404) 718–8833.

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.

Elaine L. Baker,

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

[FR Doc. 2015–13315 Filed 6–1–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.598]

Announcement of the Award of a Single-Source Program Expansion Supplement Grant to the U.S. Committee for Refugees and Immigrants in Arlington, VA

AGENCY: Office of Refugee Resettlement, ACF, HHS.

ACTION: Announcement of the award of a single-source program expansion supplement to the U.S. Committee for Refugees and Immigrants (USCRI) to support expanded services to foreign trafficking victims, potential trafficking victims, and certain family members.

SUMMARY: The Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR) announces the award of a single-source program expansion supplement grant to U.S. Committee for Refugees and Immigrants in Arlington, Virginia for a total of $1,060,805.

The supplemental funding will ensure that clients' essential needs, such as housing, transportation, communication, food, and medical care, will be met. The supplemental funding will also ensure that USCRI has sufficient staff for the increased number of cases.

DATES: The period of support under these supplements is June 01, 2015 through September 29, 2015.

FOR FURTHER INFORMATION CONTACT: Maggie Wynne, Director, Division of Anti-Trafficking in Persons, Office of Refugee Resettlement, 901 D Street SW., Washington, DC 20024, Telephone (202) 401–4664. Email: *maggie.wynne@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: The National Human Trafficking Victim Assistance Program (NHTVAP) provides funding for comprehensive case management services to victims of trafficking and certain family members on a per capita basis. The NHTVAP grantees help clients gain access to housing, employability services, mental health screening and therapy, medical care, and some legal services. During FY 2015, a grantee, U.S. Committee for Refugees and Immigrants (USCRI), served more clients than it had planned for in its budget for the year. Without the additional funding, USCRI would have to make significant cuts in services to current clients and limit the enrollment of new clients. Also, without additional funding USCRI would not have sufficient programmatic support for the increase in client enrollments. With the supplemental funding, USCRI will be able to ensure that all of the clients' essential needs will be met.

Statutory Authority: Trafficking Victims Protection Act of 2000 (TVPA), as amended, Section 107(b)(1)(B), 22 U.S.C. 7105(b)(1)(B), authorizes funding for benefits and services to foreign victims of severe forms of trafficking in persons in the United States, potential victims of trafficking seeking HHS Certification, and certain family members.

Christopher Beach,

Senior Grants Policy Specialist, Office of Administration.

[FR Doc. 2015–13177 Filed 6–1–15; 8:45 am]

BILLING CODE 4184–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0279]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Prescription Drug Marketing Act of 1987; Policies, Requirements, and Administrative Procedures

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing