Agenda items are subject to change as priorities dictate. An agenda is also posted on the NIOSH Web site (*http://www.cdc.gov/niosh/bsc/*).

Contact Person For More Information: John Decker, Executive Secretary, BSC, NIOSH, CDC, 1600 Clifton Road NE., MS–E20, Atlanta, Georgia 30333, Telephone: (404) 498–2500, Fax: (404) 498–2526.

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.

Gary Johnson,

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

[FR Doc. 2014–18687 Filed 8–6–14; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Subcommittee on Procedures Review, Advisory Board on Radiation and Worker Health (ABRWH or 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: 11:00 a.m.–5:00 p.m., Eastern Time, August 28, 2014.

Place: Audio Conference Call via FTS Conferencing.

Status: Open to the public. 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, 1–866–659–0537 and the passcode is 9933701.

Background: The ABRWH 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 compensation program. Key functions of the ABRWH 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 ABRWH is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 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, providing advice to 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 a reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee on Procedures Review was established to aid the ABRWH in carrying out its duty to advise the Secretary, HHS, on dose reconstructions. The Subcommittee on Procedures Review is responsible for overseeing, tracking, and participating in the reviews of all procedures used in the dose reconstruction process by the NIOSH Division of Compensation Analysis and Support (DCAS) and its dose reconstruction contractor (Oak Ridge Associated Universities—ORAU).

Matters for Discussion: The agenda for the Subcommittee meeting includes: discussion of procedures in the following ORAU and DCAS technical documents: ORAU Team Technical Information Bulletin (OTIB) 0034 (''Internal Dose Coworker Data for X–10''), OTIB 0054 (''Fission and Activation Product Assignment for Internal Dose-Related Gross Beta and Gross Gamma Analyses''), OTIB 0083 (''Dissolution Models for Insoluble Plutonium 238''), Program Evaluation Report (PER) 011 (''K–25 [Technical Basis Document] TBD and TIB Revisions''), PER 018 (''Los Alamos National Laboratory TBD Revision, Rev. 00,''), PER 031 (''Y–12 TBD Revisions''), PER 033 (''Reduction Pilot Plant TBD Revision''), PER 038 (''Hooker Electrochemical TBD Revisions''); Update on Review of ORAU Team Report 0053 (''Stratified Co-Worker Sets''); and a continuation of the comment-resolution process for other dose reconstruction procedures under review by the Subcommittee.

The agenda is subject to change as priorities dictate.

Contact Person for More Information: Theodore Katz, Designated Federal Officer, NIOSH, CDC, 1600 Clifton Road NE., Mailstop E–20, Atlanta, Georgia 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.

Gary Johnson,

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

[FR Doc. 2014–18686 Filed 8–6–14; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.568]

Reallotment of FY 2013 Funds for the Low Income Home Energy Assistance Program (LIHEAP)

AGENCY: Office of Community Services, ACF, HHS.

ACTION: Notice of determination concerning Federal Fiscal Year (FFY) 2013 funds available for reallotment.

SUMMARY: The Administration for Children and Families (ACF), Office of Community Services (OCS), Division of Energy Assistance (DEA) announces the reallotment of $10,880,543 of FFY 2013 funds for the Low Income Home Energy Assistance Program (LIHEAP).

FOR FURTHER INFORMATION CONTACT: Lauren Christopher, Director, Division of Energy Assistance, Office of Community Services, 370 L'Enfant Promenade SW., Washington, DC 20447 Telephone (202) 401–4870; email: *lauren.christopher@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: In accordance with Section 2607(b)(1) of the Low Income Home Energy Assistance Act (the Act), Title XXVI of the Omnibus Budget Reconciliation Act of 1981 (42 U.S.C. 8621, *et seq.*), as amended, a notice was published in the **Federal Register** on January 14, 2014 announcing the Secretary's preliminary determination that $2,192,230 of FFY 2013 funds for the Low Income Home Energy Assistance Program (LIHEAP) may be available for reallotment. Subsequent to the publication of this notice, two additional grantees reported $8,688,313 of funds for reallotment. Thus, a total of $10,880,543 was reported by grantees as available for reallotment from FY 2013.

These funds became available from the following grantees:

REALLOTMENT AMOUNTS OF FFY 2013 LIHEAP FUNDS

Grantee name	FY 2013 Reallotment amount
State of Nebraska	$2,180,356.00
State of South Carolina	7,358,414.00
State of Utah	1,329,899.00
Delaware Tribe of Indians	9,793.00
Salt River Pima-Maricopa Indian Community	2,081.00
Total	10,880,543.00

Pursuant to the statute cited above, these funds were reallotted on June 17, 2014 to all current LIHEAP grantees by distributing the total reallotted funds under the formula Congress set for FFY 2014 funding. The only exception is that grantees whose allocations would have been less than $25 did not receive an award.

The reallotted funds may be used for any purpose authorized under LIHEAP. Grantees must add these funds to their total LIHEAP funds payable for FFY 2014 for purposes of calculating statutory caps on administrative costs, carryover, assurance 16 activities, and weatherization assistance.

Statutory Authority: 45 CFR 96.81 and 42 U.S.C. 8621 et seq.

Jeannie Chaffin,

Director, Office of Community Services.

[FR Doc. 2014–18672 Filed 8–6–14; 8:45 am]

BILLING CODE 4180–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–1104]

Agency Information Collection Activities; Proposed Collection; Comment Request; State Petitions for Exemption From Preemption

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on our proposed collection of certain information. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice invites comments on the information collection provisions of our reporting requirements contained in existing FDA regulations governing state petitions for exemption from preemption.

DATES: Submit either electronic or written comments on the collection of information by October 6, 2014.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, we are publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, we invite comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of our 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 respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

State Petitions for Exemption From Preemption—21 CFR 100.1(d) (OMB Control No. 0910–0277)—Extension

Under section 403A(b) of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 343–1(b)), states may petition FDA for exemption from Federal preemption of state food labeling and standard of identity requirements. Section 100.1(d) (21 CFR 100.1(d)) sets forth the information a state is required to submit in such a petition. The information required under § 100.1(d) enables FDA to determine whether the state food labeling or standard of identity requirement satisfies the criteria of section 403A(b) of the FD&C Act for granting exemption from Federal preemption.

We estimate the burden of this collection of information as follows: