in, an entity is required to provide their Data Universal Numbering System (DUNS) number. FSRS then pulls core data about the entity from their System for Award Management (SAM) registration to include the legal business name, physical address, mailing address and Commercial and Government Entity (CAGE) code. The entity completes the FSRS registration by providing contact information within the entity for approval.

If a prime awardee has already registered in FSRS to report contracts-related Transparency Act financial data, a new log-in will not be required. In addition, if a prime awardee had a user account in the Electronic Subcontract Reporting System (eSRS), a new log-in will not be required.

B. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FSRS Registration Requirements for Prime Grant Awardees, whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology. A 60-day notice, requesting comments was published in the **Federal Register** at 78 FR 79454 on December 30, 2013, no comments were received.

C. Annual Reporting Burden

Respondents: 1,844.
Responses Per Respondent: 1.
Total annual responses: 1,844.
Hours Per Response: .5.
Total Burden Hours: 922.
Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 3090–0291, FSRS Registration Requirements for Prime Grant Awardees, in all correspondence.

Dated: March 18, 2014.

Sonny Hashmi,

Deputy Chief Information Officer, Office of the Deputy CIO.

[FR Doc. 2014–06555 Filed 3–24–14; 8:45 am]

BILLING CODE 6820–XY–P

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) announce the following meeting for the aforementioned committee:

Times and Dates: 9:00 a.m.–5:00 p.m., April 10, 2014.
9:00 a.m.–12:00 p.m., April 11, 2014.
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 *www.cdc.gov/hicpac.*
Purpose: The Committee is charged with providing advice and guidance to the Director, Division of Healthcare Quality Promotion, the Director, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), the Director, CDC, the Secretary, Health and Human Services regarding (1) the practice of healthcare infection prevention and control; (2) strategies for surveillance, prevention, and control of infections, antimicrobial resistance, and related events in settings where healthcare is provided; and (3) periodic updating of CDC guidelines and other policy statements regarding prevention of healthcare-associated infections and healthcare-related conditions.
Matters for Discussion: The agenda will include updates on the Draft Guideline to Prevent Surgical Site Infections, CDC and Division of Healthcare Quality Promotion's (DHQP's) activities for prevention and surveillance of healthcare associated infections (HAI), core infection prevention and control practices and the Draft Health Care Personnel guideline.
Agenda items are subject to change as priorities dictate.
Contact Person for More Information: Erin Stone, M.S., HICPAC, Division of Healthcare Quality Promotion, NCEZID, CDC, 1600 Clifton Road, NE., Mailstop A–07, Atlanta, Georgia 30333 Telephone (404) 639–4045. Email: *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.

Elaine L. Baker,

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

[FR Doc. 2014–06455 Filed 3–24–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention (CDC)

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., EST, April 16, 2014.
Place: Audio Conference Call via FTS Conferencing. The USA toll-free, dial-in number is 1–866–659–0537 and the pass code is 9933701.
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 020 (''Blockson TBD Revision''), 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''); discussion of estimating radiation doses associated with localized skin exposures to uranium at Atomic Weapons Employer facilities; 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.

Elaine L. Baker,

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

[FR Doc. 2014–06452 Filed 3–24–14; 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 Research Approaches to Improve the Care and Outcomes of People Living with Spina Bifida, FOA DD14–002, 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:

Times and Dates: 9:00 a.m.–6:00 p.m., April 14, 2014 (Closed).

9:00 a.m.–6:00 p.m., April 15, 2014 (Closed).

9:00 a.m.–6:00 p.m., April 16, 2014 (Closed).

9:00 a.m.–6:00 p.m., April 17, 2014 (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 ''Research Approaches to Improve the Care and Outcomes of People Living with Spina Bifida, FOA DD14–002, initial review.''

Contact Person For More Information: M. Chris Langub, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway NE., Mailstop F–80, Atlanta, Georgia 30341, Telephone: (770) 488–3585, *EEO6@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. 2014–06450 Filed 3–24–14; 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 A Creutzfeldt-Jakob Disease (CJD) Lookback Study: Assessing the Risk of Blood Borne Transmission of Classic Forms of Creutzfeldt-Jakob Disease, Funding Opportunity Announcement (FOA) CK14–005, 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: 2:00 p.m.–3:30 p.m., EST, April 15, 2014 (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 ''A Creutzfeldt-Jakob Disease (CJD) Lookback Study: Assessing the Risk of Blood Borne Transmission of Classic Forms of Creutzfeldt-Jakob Disease, FOA CK14–005, initial review.''

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. 2014–06451 Filed 3–24–14; 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 Grants for Injury Control