Time And Date: 8:30 a.m.–5 p.m., July 11, 2007 (Closed).

Place: CDC Roybal Campus, 1600 Clifton Road, Bldg. 19, Conference Room 232, Auditorium B2, Atlanta, GA 30333, Telephone (404) 639–8838.

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 review, discussion, and evaluation of the ''DGA International Laboratory Branch Review Panel, and the Extramural Review of Intramural Operational Research.''

Contact Person for More Information: Deborah Birx, Global AIDS Program, Director, CDC, Corporate Square, Bldg. 1, Room 1506, Mail Stop E–04, Atlanta, GA 30329, Telephone (404) 639–6137.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: June 6, 2007.

Elaine L. Baker,

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

[FR Doc. E7–11280 Filed 6–11–07; 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; Occupational Safety and Health Research, Program Announcement (PA) 07–318, and Exploratory Developmental Grants, PA PAR–06–552

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: 9 a.m.–5 p.m., July 11, 2007 (Closed).

Place: Marriott Waterfront, 700 Aliceanna Street, Baltimore, MD 21202.

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 review, discussion, and evaluation of research grant applications in response to PA 07–318, ''Occupational Safety and Health Research,'' and PAR 06–552, ''Exploratory Developmental Grants.''

Contact Person for More Information: Stephen Olenchock, Ph.D., Scientific Review Administrator, 1095 Willowdale Road, Morgantown, WV 26506, telephone 304.285.6271.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: June 6, 2007.

Elaine L. Baker,

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

[FR Doc. E7–11288 Filed 6–11–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health (NIOSH), Safety and Occupational Health Study Section (SOHSS)

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:

Times and Dates:

8 a.m.–5 p.m., June 28, 2007 (Closed).

8 a.m.–5 p.m., June 29, 2007 (Closed).

Place: Embassy Suites Hotel, 1900 Diagonal Road, Alexandria, Virginia 22314, telephone 703–684–5900, fax 703–684–1403.

Purpose: The Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) received in response to the Institute's standard grants review and funding cycles pertaining to research issues in occupational safety and health, and allied areas.

It is the intent of NIOSH to support broad-based research endeavors in keeping with the Institute's program goals. This will lead to improved understanding and appreciation for the magnitude of the aggregate health burden associated with occupational injuries and illnesses, as well as to support more focused research projects, which will lead to improvements in the delivery of occupational safety and health services, and the prevention of work-related injury and illness. It is anticipated that research funded will promote these program goals.

Matters To Be Discussed: The meeting will convene to address matters related to the conduct of Study Section business and for the study section to consider safety and occupational health-related grant applications. These portions of 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, Centers for Disease Control and Prevention, pursuant to Section 10(d) Public Law 92–463.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Price Connor, PhD, NIOSH Health Scientist, 1600 Clifton Road, NE., Mailstop E–20, Atlanta, Georgia 30333, telephone 404–498–2511, fax 404–498–2571.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: June 5, 2007.

Elaine L. Baker,

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

[FR Doc. E7–11281 Filed 6–11–07; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006D–0020]

Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Intervertebral Body Fusion Device; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of the guidance document entitled ''Class II Special Controls Guidance Document: Intervertebral Body Fusion Device.'' It was developed as a special control to support the reclassification of intervertebral body fusion devices that contain bone grafting material from class III (premarket approval) into class II (special controls). The guidance document describes a means by which these intervertebral body fusion devices may comply with the requirement of special controls for class II devices. Elsewhere in this issue of the **Federal Register**, FDA is publishing a final rule to reclassify the intervertebral body fusion device that contain bone grafting material from class III into class II (special controls) and retain those that contain any therapeutic biologic (e.g., bone morphogenic protein) in class III.

DATES: Submit written or electronic comments on this guidance at any time.