Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

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

[FR Doc. E7–2080 Filed 2–7–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Health Statistics: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Board of Scientific Counselors, National Center for Health Statistics, Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through January 19, 2009.

For information, contact Virginia Cain, Ph.D., Executive Secretary, Board of Scientific Counselors, National Center for Health Statistics, Centers for Disease Control and Prevention, Department of Health and Human Services, Metro IV Building, 3311 Toledo Road, Hyattsville, Maryland 20782, telephone 301–458–4395 or fax 301–458–4020.

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.

Dated: February 2, 2007.

Elaine L. Baker,

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

[FR Doc. E7–2076 Filed 2–7–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Task Force on Community Preventive Services

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:

Name: Task Force on Community Preventive Services.

Times and Dates: 8 a.m.–5:15 p.m. EST, February 14, 2007. 8 a.m.–12:30 p.m. EST, February 15, 2007.

Place: Centers for Disease Control and Prevention, 2500 Century Parkway, Atlanta, GA 30329.

Status: Open to the public, limited only by the space available.

Purpose: The mission of the Task Force is to develop and publish the *Guide to Community Preventive Services (Community Guide),* which is based on the best available scientific evidence and current expertise regarding essential public health and what works in the delivery of those services.

Matters to be discussed: Agenda items include: controlling obesity; worksite health promotion and the assessment of health risks with feedback; alcohol outlet density; asthma; updating existing *Community Guide* reviews; and dissemination activities and projects in which the *Community Guide* is used.

Agenda items are subject to change as priorities dictate.

Persons interested in reserving a space for this meeting should call Tony Pearson-Clarke at 404–498–0972 by close of business on February 9, 2007.

Contact person or additional information: Tony Pearson-Clarke, Community Guide Branch, Coordinating Center for Health Information and Service, National Center for Health Marking, Division of Health Communication and Marketing, 1600 Clifton Road, M/S E–69, Atlanta, GA 30333, phone: 404–498–0972.

Dated: January 31, 2007.

James D. Seligman,

Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E7–2078 Filed 2–7–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Prospective Grant of Co-Exclusive License

AGENCY: Technology Transfer Office, Centers for Disease Control and Prevention (CDC), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: This is a notice in accordance with 35 U.S.C. 209(e) and 37 CFR 404.7(a)(1)(i) that the Centers for Disease Control and Prevention (CDC), Technology Transfer Office, Department of Health and Human Services (DHHS), is contemplating the grant of a limited field of use, exclusive license in China, and a co-exclusive worldwide (excluding China) license to practice the invention embodied in the patent application referred to below to Ringpu (Baoding) Biologics and Pharmaceuticals Co. LTD., having a place of business in Baoding City, Hebel Province, PR China. CDC intends to grant rights to practice this invention (in territories other than China) to no more than two other co-licensees. The patent rights in these inventions have been assigned to the government of the United States of America. The patent application to be licensed is:

Provisional Patent Application

Title: Method of Sequencing Whole Viral Genomes, Related Compositions, and Genome Sequences.

Serial No. 60/727,038.

Filing date: 10/14/2005.

PCT Patent Application

Title: Rabies Virus Compositions and Methods.

Serial No.: N/A.

Filing Date: 10/13/2006.

Domestic Status: N/A.

Issue Date: patent pending.

The prospective exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7.

The critical feature of this technology is the ERA rabies virus whole genome DNA sequence. With the availability of the entire rabies genome, a recombinant vaccine can be developed using reverse genetics. The vaccines that can be developed using this genome are fundamentally different from classic ones that are being produced. The technology is being applied to other negative stranded RNA viruses.

ADDRESSES: Requests for a copy of these patent applications, inquiries, comments, and other materials relating to the contemplated license should be directed to Andrew Watkins, Director, Technology Transfer Office, Centers for Disease Control and Prevention (CDC), 4770 Buford Highway, Mailstop K–79, Atlanta, GA 30341, *telephone:* (770) 488–8610; *facsimile:* (770) 488–8615. Applications for an exclusive license to the territory of China filed in response to this notice will be treated as objections to the grant of the contemplated exclusive license. Only written comments and/or applications for a license which are received by CDC within thirty days of this notice will be

considered. Comments and objections submitted in response to this notice will not be made available for public inspection, and, to the extent permitted by law, will not be released under the Freedom of Information Act, 5 U.S.C. 552. A signed Confidential Disclosure Agreement will be required to receive a copy of any pending patent application.

Dated: January 31, 2007.

James D. Seligman,

Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E7–2077 Filed 2–7–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Circulatory System Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on March 1, 2007, from 8 a.m. to 5:30 p.m., and March 2, 2007, from 8 a.m. to 6 p.m.

Location: Hilton Washington DC North/Gaithersburg, Salons A, B, and C, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: James Swink, Center for Devices and Radiological Health (HFZ–450), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 240–276–4179, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512625. Please call the Information Line for up-to-date information on this meeting.

Agenda: On March 1, 2007, the committee will discuss and make recommendations regarding the premarket approval application, sponsored by Medtronic Inc., for the Chronicle Implantable Hemodynamic Monitoring System. This implantable device is intended to reduce hospitalization events or equivalent events for worsening heart failure in patients with moderate to advanced heart failure. On March 2, 2007, the committee will discuss and make recommendations regarding clinical trial designs for Patent Foreman Ovale closure devices intended to prevent recurrent stroke.

FDA intends to make background material available to the public no later than 1 business day before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/ohrms/dockets/ac/acmenu.htm*, click on the year 2007 and scroll down to the appropriate advisory committee link.

Procedure: On March 1, 2007, from 8 a.m. to 5:30 p.m., and March 2, 2007, from 8 a.m. to 10 a.m. and 12 p.m. to 6 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before February 23, 2007. Oral presentations from the public will be scheduled for approximately 30 minutes at the beginning of committee deliberations on each day and for approximately 30 minutes near the end of the committee deliberations on each day. Those desiring to make formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before February 15, 2007. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by February 16, 2007.

Closed Presentation of Data: On March 2, 2007, from 10 a.m. to 12 p.m., the meeting will be closed to permit the discussion and review of trade secret and/or confidential information (5 U.S.C. 552b(c)(4)) presented by sponsors.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, at 301–827–7291, at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: February 1, 2007.

Randall W. Lutter,

Associate Commissioner for Policy and Planning.

[FR Doc. E7–2122 Filed 2–7–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005D–0091]

Guidance for Industry on User Fee Waivers for Fixed Dose Combination and Co-Packaged Human Immunodeficiency Virus Drugs for the President's Emergency Plan for Acquired Immunodeficiency Syndrome Relief; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a guidance for industry entitled ''User Fee Waivers for FDC and Co-Packaged HIV Drugs for PEPFAR.'' This guidance describes the circumstances under which user fees will not be assessed for certain applications for fixed dose combination (FDC) and co-packaged versions of previously approved antiretroviral therapies for the treatment of human immunodeficiency virus (HIV) under the President's Emergency Plan for Acquired Immunodeficiency Syndrome Relief (PEPFAR). The guidance also describes some circumstances under which most of the applications that will be assessed fees may be eligible for a public health or a barrier-to-innovation waiver.

DATES: Submit written or electronic comments on agency guidances at any time.

ADDRESSES: Submit written requests for single copies of the guidance to the Division of Drug Information (HFD–