Survey of Consumer Finance (SCF) will be the latest in a triennial series, which began in 1983, that provides comprehensive data for U.S. families on the distribution of assets and debts, along with related information and other data items necessary for analyzing financial behavior. These are the only surveys conducted in the United States that provide such financial data for a representative sample of households. Data for the SCF are collected by interviewers using a computer program. While some questions may be deleted and others modified, only minimal changes will be made to the questionnaire in order to preserve the time series properties of the data. The pretest would be conducted during 2006 and the survey would be conducted between May 2007 and January 2008.

Board of Governors of the Federal Reserve System, July 5, 2006.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. E6–10782 Filed 7–10–06; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSTEM

Government in the Sunshine Act; Meeting Notice

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 11:30 a.m., Monday, July 17, 2006.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, NW., Washington, DC 20551

STATUS: Closed.

MATTERS TO BE CONSIDERED: 1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any items carried forward from a previously announced meeting.

FOR MORE INFORMATION PLEASE CONTACT: Michelle Smith, Director, or Dave Skidmore, Assistant to the Board, Office of Board Members at 202–452–2955.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board's Web site at *http://www.federalreserve.gov* for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: July 7, 2006.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 06–6177 Filed 7–7–06; 3:57 pm]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator; American Health Information Community Biosurveillance Data Steering Group Meeting

ACTION: Announcement of meeting.

SUMMARY: This notice announces the first meeting of the American Health Information Community Biosurveillance Data Steering Group in accordance with the Federal Advisory Committee Act (Pub. L. No. 92–463, 5 U.S.C., App.).

DATES: July 7, 2006 from 10 a..m. to 2 p.m.

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090.

FOR FURTHER INFORMATION CONTACT: *http://www.hhs.gov/healthit/ahic.html.*

SUPPLEMENTARY INFORMATION: The Biosurveillance Data Steering Group must convene in early July 2006 in advance of the final deliverable from the Health Information Technology Standards Panel related to the Biosurveillance Use Case.

The meeting will be available via internet access. Go to *http://www.hs.gov/healthit/ahic.html* for additional information on the meeting.

Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator.

[FR Doc. 06–6113 Filed 7–10–06; 8:45 am]

BILLING CODE 4150–24–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Meeting of the National Advisory Council for Healthcare Research and Quality

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Notice of public meeting.

SUMMARY: In accordance with section 10(a) of the Federal Advisory Committee Act, this notice announces a meeting of the National Advisory Council for Healthcare Research and Quality.

DATES: The meeting will held on Friday, July 21, 2006, from 8:30 a.m. to 4 p.m. and is open to the public.

ADDRESSES: The meeting will held in The Eisenberg Conference Center, the Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850. The public is reminded to bring a photo ID to enter a Federal building.

FOR FURTHER INFORMATION CONTACT: Deborah Queenan, Coordinator of the Advisory Council, at the Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850, (301) 427–1330. For press-related information, please contract Karen Migdail at (301) 427–1855.

If sign language interpretation or other reasonable accommodation for a disability is needed, please contact Mr. Donald L. Inniss, Director, Office of Equal Employment Opportunity Program, Program Support Center, on (301) 443–1144 no later than July 14, 2006. Agenda, roster, and minutes from previous council meetings are available from Ms. Bonnie Campbell, Committee Management Officer, Agency for Healthcare Quality and Research, 540 Gaither Road, Rockville, Maryland 20850. Ms. Campbell's phone number is (301) 427–1554.

SUPPLEMENTARY INFORMATION:

I. Purpose

Section 931 of the Public Health Service Act (42 U.S.C. 299c) established the National Advisory Council for Healthcare Research and Quality. In accordance with its statutory mandate, the Council is to advise the Secretary of the Department of Health and Human Services and the Director, Agency for Healthcare Research and Quality (AHRQ), on matters related to actions of the Agency to enhance the quality, improve the outcomes, reduce the costs of health care services, improve access to such services through scientific research, and to promote improvements in clinical practice and in the organization, financing and delivery of health care services.

The Council is composed of members of the public appointed by the Secretary, and Federal ex-officio members.

II. Agenda

On Friday, July 21, 2006, the meeting will convene at 8:30 a.m. with the call to order by the Council Chair. The agenda will include the Director's update on the status of the Agency's current research, programs, and initiatives; a presentation and discussion on AHRQ's research on

health care efficiency; and an overview of the National Healthcare Quality and Disparities Reports. The final agenda will be available on AHRQ's Web site at *http://www.ahrq.gov* no later than July 14, 2006.

The meeting will adjourn at 4 p.m.

Dated: July 5, 2006.

Carolyn M. Clancy,

Director.

[FR Doc. 06–6164 Filed 7–7–06; 2:05 pm]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and ControlSpecial Emphasis Panel: Targeted Evaluation of the President's Emergency Plan for AIDS Relief (PEPFAR) Funded Prevention of Mother-to-Child HIV Transmission (PMTCT), and Adherence to Antiretroviral Therapy (ART) Programs, Contract Solicitation Numbers (CSN) 2006–N–08428, 2006–N–08429, and 2006–N–08430

Correction: This notice was published in the **Federal Register** on June 9, 2006, Volume 71, Number 111, page 33456. The location of the meeting was changed due to insufficient meeting space at the Renaissance Concourse Hotel—Marriott, One Hartsfield Center Parkway, Atlanta, GA 30354. The meeting was held at the Hilton Atlanta Airport, 1031 Virginia Avenue, Atlanta, Georgia 30354.

Titles: Targeted Evaluation of the President's Emergency Plan for AIDS Relief (PEPFAR) Funded Prevention of Mother-to-Child HIV Transmission (PMTCT), and Adherence to Antiretroviral Therapy (ART) Programs, Contract Solicitation Numbers (CSN) 2006–N–08428, 2006–N–08429, and 2006–N–08430.

For Further Information Contact: Amy L. Sandul, Health Scientist, National Center for HIV, STD, and Tuberculosis Prevention, Centers for Disease Control and Prevention, 1600 Clifton Road, NE., MS E–41, Atlanta, GA 30333, Telephone 404–639–6485.

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: July 3, 2006.

Alvin Hall,

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

[FR Doc. E6–10774 Filed 7–10–06; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005E–0236]

Determination of Regulatory Review Period for Purposes of Patent Extension; MULTIHANCE

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for MULTIHANCE and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of Patents and Trademarks, Department of Commerce, for the extension of a patent that claims that human drug product.

ADDRESSES: Submit written comments and petitions to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments*.

FOR FURTHER INFORMATION CONTACT: Beverly Friedman, Office of Regulatory Policy (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: The Drug Price Competition and Patent Term Restoration Act of 1984 (Public Law 98–417) and the Generic Animal Drug and Patent Term Restoration Act (Public Law 100–670) generally provide that a patent may be extended for a period of up to 5 years so long as the patented item (human drug product, animal drug product, medical device, food additive, or color additive) was subject to regulatory review by FDA before the item was marketed. Under these acts, a product's regulatory review period forms the basis for determining the amount of extension an applicant may receive.

A regulatory review period consists of two periods of time: A testing phase and an approval phase. For human drug products, the testing phase begins when the exemption to permit the clinical investigations of the human drug product becomes effective and runs until the approval phase begins. The approval phase starts with the initial submission of an application to market the human drug product and continues until FDA grants permission to market the product. Although only a portion of a regulatory review period may count toward the actual amount of extension that the Director of Patents and Trademarks may award (for example, half the testing phase must be subtracted, as well as any time that may have occurred before the patent was issued), FDA's determination of the length of a regulatory review period for a human drug product will include all of the testing phase and approval phase as specified in 35 U.S.C. 156(g)(1)(B).

FDA recently approved for marketing the human drug product MULTIHANCE (gadobenate dimeglumine). MULTIHANCE is indicated for intravenous use in magnetic resonance imaging (MRI) of the central nervous system in adults to visualize lesions with abnormal blood brain barrier or abnormal vascularity of the brain, spine, and associated tissues. Subsequent to this approval, the Patent and Trademark Office received a patent term restoration application for MULTIHANCE (U.S. Patent No. 4,916,246) from Bracco International B.V., and the Patent and Trademark Office requested FDA's assistance in determining this patent's eligibility for patent term restoration. In a letter dated July 8, 2005, FDA advised the Patent and Trademark Office that this human drug product had undergone a regulatory review period and that the approval of MULTIHANCE represented the first permitted commercial marketing or use of the product. Thereafter, the Patent and Trademark Office requested that FDA determine the product's regulatory review period.

FDA has determined that the applicable regulatory review period for MULTIHANCE is 3,789 days. Of this time, 2,482 days occurred during the testing phase of the regulatory review period, while 1,307 days occurred during the approval phase. These periods of time were derived from the following dates:

1. *The date an exemption under section 505(i) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 355) became effective*: July 12, 1994. FDA has verified the applicant's claim that the date the investigational new drug application became effective was on July 12, 1994.

2. *The date the application was initially submitted with respect to the human drug product under section*