N.V.,all of Brussels, Belgium, Fortis Bank Nederland (Holding) N.V., Utrecht, Netherlands, and RFS Holdings B.V., Amsterdam, Netherlands, is revised to read as follows:

A. Federal Reserve Bank of Boston (Richard Walker, Community Affairs Officer) P.O. Box 55882, Boston, Massachusetts 02106-2204:

1. Royal Bank of Scotland Group, plc, Edinburgh, Scotland, Banco Santander Central Hispano, S.A., Madrid, Spain, Santander Holanda B.V., Delft, Netherlands, Fortis N.V., Utrecht, Netherlands, Fortis S.A./N.V.,Fortis Brussels, S.A./N.V., Fortis Bank, all of Brussels, Belgium, Fortis Bank Nederland (Holding) N.V., Utrecht, Netherlands, and RFS Holdings B.V., Amsterdam, Netherlands; to control ABN AMRO Holding N.V. Amsterdam, Netherlands, and thereby indirectly acquire ABN AMRO North American Holding Company, LaSalle Bank Corporation, LaSalle Bank National Association, all of Chicago, Illinois, and LaSalle Bank Midwest National Association, Troy, Michigan. In connection with this proposal Fortis Bank Nederland (Holding) N.V., Santander Holand B.V. and RFS Holdings B.V. have applied to become bank holing companies.

In addition, each of The Royal Bank of Scotland Group, plc, The Royal Bank of Scotland plc, RBSG International Holdings Limited, all of Edinburgh, Scotland, and Citizens Financial Group, Inc., Providence, Rhode Island, has applied to acquire control of ABN AMRO North American Holding Company, LaSalle Bank Corporation, LaSalle Bank National Association, and LaSalle Bank Midwest National Association in a transfer subsequent to the acquisition of control of ABN AMRO Holding N.V.

Comments on this application must be received by July 25, 2007.

Board of Governors of the Federal Reserve System, July 9, 2007.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E7–13530 Filed 7–11–07; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Ethics Subcommittee, Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), CDC announces the following meeting for the aforementioned subcommittee.

Times and dates: 1 p.m.–5:30 p.m., August 9, 2007. 8:30 a.m.–3:30 p.m., August 10, 2007.

Place: CDC, 1825 Century Center, Conference Room 1 A/B, Atlanta, GA 30345.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 75 people. To accommodate public participation in the meeting, a conference telephone line will be available. The public is welcome to participate during the public comment periods by calling (866) 919–3560 and entering code 4168828. The public comment periods are tentatively scheduled from 4:45 p.m.–5 p.m. on August 9, 2007 and from 3 p.m.–3:15 p.m. on August 10, 2007.

Purpose: The Ethics Subcommittee will provide counsel to the ACD, CDC regarding a broad range of public health ethics questions and issues arising from programs, scientists, and practitioners.

Matters To Be Discussed: Agenda items will include: Ethical Guidance for Public Health Emergency Preparedness and Response, Ethical Issues relating to CDC Partnerships, Public Health Ethics and Genomics, Ethical Guidance for Non-Research Data Collections, and Updates on Ethical Issues relating to Pandemic Influenza Preparedness. Agenda items are subject to change as priorities dictate.

For security reasons, members of the public interested in attending the meeting should contact the person below. The deadline for notification of attendance is August 2, 2007.

Contact Person for More Information: Drue Barrett, Ph.D., Designated Federal Official, Ethics Subcommittee, CDC, 1600 Clifton Road, NE., M/S D–50, Atlanta, Georgia 30333. Telephone (404) 639–4690. E-mail: *dbarrett@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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: July 5, 2007.

Elaine L. Baker,

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

[FR Doc. E7–13523 Filed 7–11–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Antiviral Drugs 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: Antiviral Drugs 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 September 5, 2007, from 8 a.m. to 4 p.m. and on September 6, 2007, from 9 a.m. to 1 p.m.

Location: On September 5, 2007, the committee will meet at the Hilton Washington DC/Silver Spring, The Ballrooms, 8727 Colesville Rd., Silver Spring, MD. The hotel telephone number is 301–589–5200. On September 6, 2007, the committee will meet in closed session at FDA, White Oak Headquarters, rm. 2046, 10903 New Hampshire Ave., Silver Spring, MD.

Contact Person: Cicely Reese, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301–827–7001, FAX: 301–827–6776, e-mail: *Cicely.Reese@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512531. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: On September 5, 2007, the committee will discuss new drug application (NDA) 22–145, raltegravir potassium, integrase inhibitor 400 milligram tablets, Merck & Co., Inc., for the treatment of Human Immunodeficiency Virus-1 (HIV–1) infection in combination with other