Federal Communications Commission.

Marlene H. Dortch,

Secretary.

[FR Doc. 03–29346 Filed 11–20–03; 1:46 pm]

BILLING CODE 6712–01–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies; Correction

This notice corrects a notice (FR Doc. 03–28720) published on page 65070 of the issue for Tuesday, November 18, 2003.

Under the Federal Reserve Bank of Richmond heading, the entry for Bank of America Corporation, Charlotte, North Carolina, is revised to read as follows:

A. Federal Reserve Bank of Richmond (A. Linwood Gill, III, Vice President) 701 East Byrd Street, Richmond, Virginia 23261-4528:

1. Bank of America Corporation, Charlotte, North Carolina; to merge with FleetBoston Financial Corporation, Boston, Massachusetts, and thereby indirectly acquire Fleet National Bank, Providence, Rhode Island, and Fleet Maine, National Association, South Portland, Maine.

In connection with this proposal, Bank of America has applied to acquire up to 19.9 percent of FleetBoston Financial Corporation, and FleetBoston Financial Corporation has an option to acquire 19.9 percent of the voting shares of Bank of America Corporation.

Comments on this application must be received by December 15, 2003.

Board of Governors of the Federal Reserve System, November 18, 2003.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 03–29239 Filed 11–21–03; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Secretary's Advisory Committee on Human Research Protections

AGENCY: Office of the Secretary, HHS.

ACTION: Notice of meeting.

SUMMARY: Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. appendix 2), notice is hereby given of the second meeting of the Secretary's Advisory Committee on Human Research Protections (SACHRP). The meeting will be open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the contact person listed below.

DATES: The meeting will be held on Thursday, December 11 and Friday, December 12, 2003, and will convene each day from approximately 8:30 a.m. until 5 p.m. EST.

ADDRESSES: The Sheraton Four Points Hotel, 1201 K Street, NW., Washington, DC, 20005.

FOR FURTHER INFORMATION CONTACT: Bernard Schwetz, D.V.M., Ph.D., Acting Executive Secretary, SACHRP, Department of Health and Human Services (HHS), Office of Public Health and Science (OPHS), 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852, (301) 496–7005, fax: (301) 402–0527, e-mail address: *sachrp@osophs.dhhs.gov* or Catherine Slatinshek, Executive Director, SACHRP, HHS, OPHS, 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852, (301) 496–5433, fax: (301) 496–0527, e-mail address: *cslatinshek@osophs.dhhs.gov.*

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. 217a, section 222 of the Public Health Service Act, as amended, HHS established SACHRP to provide expert advice and recommendations to the Secretary of HHS and the Assistant Secretary for Health on issues and topics pertaining to or associated with the protection of human research subjects.

On December 11, SACHRP will receive and discuss preliminary reports from its three subcommittees that were created by SACHRP at its July 22, 2003 meeting to address issues related to the following three topics areas: Department of Health and Human Services (HHS) regulations and policies for research involving prisoners, HHS regulations and policies for research involving children, and the accreditation of human research protection programs by non-federal accrediting bodies. On December 12, SACHRP will hold panel discussions related to human subjects research in international settings and adverse event reporting requirements under HHS and Food and Drug Administration regulations. The committee will also discuss future tasks for 2004.

Members of the public will have the opportunity to provide comments at the meeting on December 11 and 12, 2003. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed material distributed to SACHRP members should submit materials to the Acting Executive Secretary of SACHRP (contact information listed above) prior to close of business December 2, 2003.

Information about SACHRP and the draft meeting agenda will be posted on the SACHRP Web site at: *http://ohrp.osophs.dhhs.gov/sachrp/sachrp.htm.*

Dated: November 18, 2003.

Bernard A. Schwetz,

Acting Director, Office for Human Research Protections, Acting Executive Secretary, Secretary's Advisory Committee on Human Research Protections.

[FR Doc. 03–29298 Filed 11–21–03; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–04–04]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498–1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503, or by fax to (202) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project: Restriction on Travel of Persons (OMB No. 0920–0488)—Extension—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC).

In 2000, the Food and Drug Administration (FDA) and CDC consolidated regulations related to controlling the spread of communicable diseases. FDA formerly administered the regulations contained in part 1240 of title 21, Code of Federal Regulations, which pertained to interstate control of communicable diseases. These regulations may now be found in part 70 of title 42, Code of Federal Regulations.

FDA transferred 21 CFR part 1240 to CDC. This mandate regulates the interstate travel of any person who is in the communicable period of cholera, plague, smallpox, typhus, or yellow fever, or who, having been exposed to any such disease, is in the incubation period thereof. One of the sections—

formerly 21 CFR 1240.50 and now 42 CFR 70.5 (Certain communicable diseases; special requirements)—contains a requirement for reporting certain information to the Federal government. Specifically, this regulation requires any person who is in the communicable period of cholera, plague, smallpox, typhus or yellow fever, or who, having been exposed to any such disease, is in the incubation period thereof, to apply for and receive a permit from the Surgeon General or his authorized representative in order to travel from one State or possession to another.

Control of disease transmission within the States is considered to be the province of state and local health authorities, with federal assistance being sought by those authorities on a cooperative basis without application of federal regulations. The regulations formerly administered by FDA and assumed by CDC were developed to facilitate Federal action in the event of large outbreaks requiring a coordinated effort involving several states, or in the event of inadequate local control. While it is not known whether, or to what extent situations may arise in which these regulations would be invoked, contingency planning for domestic emergency preparedness is now commonplace. Should this occur, CDC will use the reporting and record-keeping requirements contained in the regulations to carry out quarantine responsibilities as required by law. The estimated annualized burden is 3,600 hours.

Regulation/purpose	Respondent	No. of applicants	No. of responses per applicant	Average burden per response
42 CFR 70.3 Application to the State of destination for a permit to move from one State to another with a communicable disease.	Any person with a communicable disease who is seeking to travel from one State to another.	2,000	1	15/60
	Attending physician	2,000	1	15/60
42 CFR 70.3 Copy of material submitted by applicant and permit issued by State health authority under this provision.	State health authority	8	250	6/60
42 CFR 70.4 Report by the master of a vessel person in charge of a conveyance of the incidence of a communicable disease occurring while in interstate transit.	The master of a vessel or person in charge of a conveyance engaged in interstate traffic when a case or potential case of a communicable disease is identified.	1,500	1	15/60
42 CFR 70.4 Copy of material submitted to State or local health authority under this provision.	State or local health authority	20	75	6/60
42 CFR 70.5 Application for a permit to move from State to State while in the communicable period of or having been exposed to smallpox.	Any person with or in the incubation period of certain communicable diseases who is seeking to travel form one State to another.	3,750	1	15/60
	Attending physician	3,750	1	15/60

Dated: November 17, 2003.

Laura Yerdon Martin,

Acting Director, Executive Secretariat, Centers for Disease Control and Prevention.

[FR Doc. 03–29213 Filed 11–21–03; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–02–04]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498–1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503, or by fax to (202) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project: Evaluation of Customer Satisfaction with the Agency for Toxic Substances and Disease Registry Internet Home Page and Links (OMB No. 0923–0028)—Reinstatement—Agency for Toxic Substances and Disease Registry (ATSDR).

ATSDR proposes to conduct customer satisfaction research for its Internet site. Information on the site focuses on prevention of exposure and adverse human health effects and diminished quality of life associated with exposure to hazardous substances from waste sites, unplanned releases, and other sources of pollution present in the environment. The site is designed to serve the general public, persons at risk for exposure to hazardous substances, and health professionals.

Approval for a similar Customer Satisfaction Survey was requested in 2002 jointly with the Centers for Disease Control and Prevention (OMB No. 0920–0449, Expiration Date 09/30/2003). The new survey is solely for ATSDR and is significantly shorter and would require less time to complete.

This research will ensure that targeted audiences find the information easy to access, clear, informative, and useful. Specifically, the research will examine whether the information is presented in an appropriate technological format and meets the needs, wants, and preferences of visitors or ''customers'' using the Web site. Results from the previous survey were utilized to redesign the ATSDR Web site—making improvements to architecture, links, organization, and content. Results from the new survey will assist ATSDR in making more improvements to the Web site in order to better serve its customers and visitors. The estimated annualized burden is 83 hours.