Estimated Annual reporting hours: 100 hours.

Estimated average hours per response: Loan participation renewal notice, 2 hours; Acquisition notice, 6 hours; Internal corporate reorganization transactions notice, 6 hours; and Section 23A additional exemption notice, 10 hours.

Number of respondents: Loan participation renewal notice, 1; Acquisition notice, 1; Internal corporate reorganization transactions notice, 12; and Section 23A additional exemption notice, 2.

General description of report: This information collection is required to evidence compliance with sections 23A and 23B of the Federal Reserve Act (12 U.S.C. 371c and 371c-1). Confidential and proprietary information collected for the purposes of the Loan Participation Renewal notice (12 CFR 223.15(b)(4)) may be protected under the authority of section (b)(4) of FOIA [5 U.S.C. 552(b)(4)]. That section of FOIA exempts commercial or financial information deemed competitively sensitive from disclosure. Respondents who desire that the information on this notice be kept confidential in accordance with section (b)(4) can request confidential treatment under the Board's rules at 12 CFR 261.15. In addition, information that is obtained as part of an examination of a financial institution is exempt from disclosure under exemption (b)(8) of FOIA. 5 U.S.C. 552(b)(8).

Abstract: On December 12, 2002, the Federal Reserve published a **Federal Register** notice [1] adopting Regulation W (Reg W) to implement sections 23A and 23B. Reg W was effective April 1, 2003. The Board issued Reg W for several reasons. First, the regulatory framework established by the Gramm-Leach-Bliley Act [2] emphasized the importance of sections 23A and 23B as a means to protect depository institutions from losses in transactions with affiliates. Second, adoption of a comprehensive rule simplified the interpretation and application of sections 23A and 23B, ensured that the statute is consistently interpreted and applied, and minimized burden on banking organizations to the extent consistent with the statute's goals. Third, issuing a comprehensive rule allowed the public an opportunity to comment on Federal Reserve interpretations of sections 23A and 23B.

The information collection requirements associated with Regulation W comprise four notices: (1) The Loan Participation Renewal notice (12 CFR 223.15(b)(4)), which is a condition to an exemption for renewals of loan participations involving problem loans; (2) the Acquisition notice (12 CFR 223.31(d)(4)), which is a condition to an exemption for a depository institution's acquisition of an affiliate that becomes an operating subsidiary of the institution after the acquisition; (3) the Internal Corporate Reorganization Transactions notice (12 CFR 223.41(d)(2)),which is a condition to an exemption for internal corporate reorganization transactions; and (4) the Section 23A Additional Exemption notice (12 CFR 223.43(b)),which provides procedures for requesting additional exemptions from the requirements of section 23A. These notifications are event-generated and must be provided to the appropriate federal banking agency and, if applicable, the Federal Reserve Board within the time periods established by the law and regulation.

Board of Governors of the Federal Reserve System, March 8, 2012.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 2012–6074 Filed 3–13–12; 8:45 am]

BILLING CODE 6210–01–P

[1] (67 FR 76603).

[2] Public Law 106–102, 113 Stat. 1338 (1999).

FEDERAL RESERVE SYSTEM

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

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The applications will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than April 9, 2012.

A. Federal Reserve Bank of Richmond (Adam M. Drimer, Assistant Vice President) 701 East Byrd Street, Richmond, Virginia 23261–4528:

1. *First Carolina Financial Services, Inc.,* Durham, North Carolina; to become a bank holding company by acquiring at least 95 percent of the voting shares of First Carolina State Bank, Rocky Mount, North Carolina, and Pisgah Community Bank, Asheville, North Carolina.

Board of Governors of the Federal Reserve System, March 9, 2012.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2012–6147 Filed 3–13–12; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0218]

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). The meeting will be open 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.

DATES: *Date and Time:* The meeting will be held on May 10, 2012, from 8 a.m. to 5:30 p.m.

ADDRESSES: FDA is opening a docket for public comment on this meeting. The docket number is FDA–2012–N–0218. The docket will open for public comment on March 14, 2012. The docket will close on May 17, 2012. Interested persons may submit either electronic or written comments regarding this meeting. Submit electronic comments to *http://www.regulations.gov*. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All

comments received will be posted without change, including any personal information provided. It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. Comments received on or before April 26, 2012, will be provided to the committee before the meeting.

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Building 31 Conference Center, the Great Room (rm. 1503), Silver Spring, MD 20993–0002. Information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/default.htm*; under the heading ''Resources for You,'' click on ''Public Meetings at the FDA White Oak Campus.'' Please note that visitors to the White Oak Campus must enter through Building 1.

Contact Person: Yvette Waples, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001, FAX: 301–847–8533, email: *AVAC@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), and follow the prompts to the desired center or product area. 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: The committee will discuss an efficacy supplement for new drug application (NDA) 21–572, TRUVADA (emtricitabine/tenofovir disoproxil fumarate) Tablet, submitted by Gilead Sciences, Inc. The supplemental application proposes an indication for Pre-Exposure Prophylaxis (PrEP) to reduce the risk of sexually acquired HIV–1 infection.

FDA intends to make background materials available to the public no later than 2 business days 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/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. All electronic and written submissions submitted to the Docket (see the **ADDRESSES** section of this document) on or before April 26, 2012, will be provided to the committee. Oral presentations from the public will be scheduled between approximately 2 p.m. and 3 p.m. Those individuals interested in making 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 April 18, 2012. 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 April 19, 2012.

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 Yvette Waples at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

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

Dated: March 8, 2012.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2012–6115 Filed 3–13–12; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub. L. 104–13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call the HRSA Reports Clearance Officer at (301) 443–1984.

Comments are invited on: (a) The proposed collection of information for the proper performance of the functions of the agency; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Voluntary Partner Surveys To Implement Executive Order 12862 in the Health Resources and Services Administration—(OMB No. 0915–0212)—[Revision]

In response to Executive Order 12862, the Health Resources and Services Administration (HRSA) is proposing to conduct voluntary customer surveys of its partners to assess strengths and weaknesses in program services and processes. HRSA partners are typically State or local governments, health care facilities, health care consortia, health care providers, and researchers. HRSA is requesting a generic approval from OMB to conduct the partner surveys.