

Commission's Rules of Practice and Procedure, 46 CFR 502.2, as well as being mailed directly to all parties of record;

Finally, it is ordered That pursuant to the terms of Rule 61 of the Commission's Rules of Practice and Procedure, 46 CFR 502.61, the final decision of the Commission in this proceeding shall be issued by August 9, 2012.

By the Commission.

Karen V. Gregory,
Secretary.

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necessary to facilitate settlement of the Federal Reserve's agency MBS transactions. The System Open Market Account Manager and the Secretary will keep the Committee informed of ongoing developments regarding the System's balance sheet that could affect the attainment over time of the Committee's objectives of maximum employment and price stability.

By order of the Federal Open Market Committee, April 9, 2012.

William B. English,
Secretary, Federal Open Market Committee.

[FR Doc. 2012-8918 Filed 4-13-12; 8:45 am]

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B. Federal Reserve Bank of San Francisco (Kenneth Binning, Vice President, Applications and Enforcement) 101 Market Street, San Francisco, California 94105-1579:

1. *Carpenter Fund Manager GP, LLC, Carpenter Fund Management Company, LLC, Carpenter Community Bancfund, L.P., Carpenter Community BanFund—A, L.P., Carpenter Community BandFund—CA, L.P., CCFW, Inc., and Carpenter Bank Partners, Inc.*, all in Irvine, California; to acquire additional voting shares, for a total of approximately 78 percent of the voting shares, of Manhattan Bancorp, and thereby indirectly acquire additional voting shares of Bank of Manhattan, National Association, both in El Segundo, California.

Board of Governors of the Federal Reserve System, April 11, 2012.

Robert deV. Frierson,
Deputy Secretary of the Board.

[FR Doc. 2012-9032 Filed 4-13-12; 8:45 am]

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FEDERAL RESERVE SYSTEM

Federal Open Market Committee; Domestic Policy Directive of March 13, 2012

In accordance with Section 271.7(d) of its rules regarding availability of information (12 CFR part 271), there is set forth below the domestic policy directive issued by the Federal Open Market Committee at its meeting held on March 13, 2012.¹

The Federal Open Market Committee seeks monetary and financial conditions that will foster price stability and promote sustainable growth in output. To further its long-run objectives, the Committee seeks conditions in reserve markets consistent with federal funds trading in a range from 0 to ¼ percent. The Committee directs the Desk to continue the maturity extension program it began in September to purchase, by the end of June 2012, Treasury securities with remaining maturities of approximately 6 years to 30 years with a total face value of \$400 billion, and to sell Treasury securities with remaining maturities of 3 years or less with a total face value of \$400 billion. The Committee also directs the Desk to maintain its existing policies of rolling over maturing Treasury securities into new issues and of reinvesting principal payments on all agency debt and agency mortgage-backed securities in the System Open Market Account in agency mortgage-backed securities in order to maintain the total face value of domestic securities at approximately \$2.6 trillion. The Committee directs the Desk to engage in dollar roll transactions as

¹ Copies of the Minutes of the Federal Open Market Committee at its meeting held on March 13, 2012, which includes the domestic policy directive issued at the meeting, are available on the Board's Web site, www.federalreserve.gov. The minutes are also published in the Federal Reserve Bulletin and in the Board's Annual Report.

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 May 11, 2012.

A. Federal Reserve Bank of Kansas City (Dennis Denney, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198-0001:

1. *FNB Bancshares, Inc.*, Independence, Kansas; to become a bank holding company by acquiring 100 percent of the voting shares of First National Bank, Independence, Kansas.

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Formations of, Acquisitions by, and Mergers of Bank Holding Companies; Correction

This notice corrects a notice (FR Doc. 2012-08198) published on page 20635 of the issue for Thursday, April 5, 2012.

Under the Federal Reserve Bank of Kansas City heading, the entry for *Arthur L. Loomis, II, Patricia A. Loomis, Genevieve E. Loomis, and Julia P. Loomis, all of Niskayuna, New York; Frederick S. Loomis, Anne M. Loomis, and J. Porter Loomis, all of Pratt, Kansas; Howard K. Loomis, Jr., Karen P. Loomis, Katherine P. Loomis, Margaret P. Loomis, and Victoria K. Loomis, all of Los Gatos, California, as individuals and/or trustees of the 2011 Arthur L. Loomis, II Gift Trust, Julia P. Loomis Revocable Trust, Arthur L. Loomis, II Revocable Trust, Genevieve E. Loomis Revocable Trust, all of Niskayuna, New York; Howard K. Loomis Revocable Trust, 2010 Howard K. Loomis Irrevocable Family Trust, Porter Legacy Trust, Florence Porter Loomis Trust, 2010 Florence Porter Loomis Irrevocable Family Trust, 2011 Frederick S. Loomis Gift Trust, 2011 J. Porter Loomis Gift Trust, all of Pratt, Kansas; 2011 Howard K. Loomis Jr. Gift Trust, The Loomis 1993 Revocable Trust, both of Los Gatos, California; and Flopper, L.P., How-Kan, L.P., and Driftwood, LLC, all of Pratt, Kansas; and all as members of the Loomis Family Group, is revised to read as follows:*

A. Federal Reserve Bank of Kansas City (Dennis Denney, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198-0001:

1. *Arthur L. Loomis, II, Patricia A. Loomis, Genevieve E. Loomis, and Julia P. Loomis, all of Niskayuna, New York; Florence Porter Loomis, Frederick S. Loomis, Anne M. Loomis, and J. Porter Loomis, all of Pratt, Kansas; Howard K. Loomis, Jr., Karen P. Loomis, Katherine P. Loomis, Margaret P. Loomis, and Victoria K. Loomis, all of Los Gatos, California, as individuals and/or trustees of the 2011 Arthur L. Loomis, II Gift Trust, the Julia P. Loomis Revocable Trust, the Arthur L. Loomis, II Revocable Trust, the Genevieve E. Loomis Revocable Trust, all of Niskayuna, New York; the Howard K. Loomis Revocable Trust, the 2010 Howard K. Loomis Irrevocable Family Trust, the Porter Legacy Trust, Florence Porter Loomis Trust, the 2010 Florence Porter Loomis Irrevocable Family Trust, the 2011 Frederick S. Loomis Gift Trust, the 2011 J. Porter Loomis Gift Trust, all of Pratt, Kansas; the 2011 Howard K. Loomis Jr. Gift Trust, The Loomis 1993 Revocable Trust, both of Los Gatos, California; and Flopper, L.P., How-Kan, L.P., and Driftwood, LLC, all of Pratt, Kansas; and all as members of the Loomis Family Group, to retain control of Krey Co. Ltd., and thereby indirectly retain control of The Peoples Bank, both in Pratt, Kansas.*

Comments on this application must be received by April 20, 2012.

Board of Governors of the Federal Reserve System, April 11, 2012.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2012-9033 Filed 4-13-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Information Program on Clinical Trials: Maintaining a Registry and Results Databank

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Library of Medicine (NLM), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on February 9, 2012 (Vol. 77, No. 27, p. 6808) and allowed 60-days for public comment. A single public

comment was received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: Information Program on Clinical Trials: Maintaining a Registry and Results Databank; **Type of Information Collection Request:** Revision of currently approved collection [OMB No. 0925-0586, expiration date 04/30/2012], **Form Number:** NA; **Need and Use of Information Collection:** The National Institutes of Health operates ClinicalTrials.gov, which was established as a clinical trial registry under section 113 of the Food and Drug Administration Modernization Act of 1997 (Pub. L. 105-115) and was expanded to include a results data bank by Title VIII of the Food and Drug Administration Amendments Act of 2007 (FDAAA). ClinicalTrials.gov collects registration and results information for clinical trials and other types of clinical studies (e.g., observational studies and patient registries) with the objectives of enhancing patient enrollment and providing a mechanism for tracking subsequent progress of clinical studies, to the benefit of public health. It is widely used by patients, physicians, and medical researchers; in particular those involved in clinical research. While many clinical studies are registered voluntarily, FDAAA requires the registration of certain applicable clinical trials of drugs and devices and the submission of results information for completed applicable clinical trials of drugs and devices that are approved, licensed, or cleared by the Food and Drug Administration. Beginning in 2009, results information was required to include information about serious and frequent adverse events. As the existing PRA clearance for this information collection nears expiration, we are making a limited number of revisions to include additional data elements that may be voluntarily submitted to describe and aid in the interpretation of any submitted adverse event information, to facilitate the registration of patient registries, and to account for the burden of establishing an account with the ClinicalTrials.gov Protocol Registration System (PRS). **Frequency of Response:** For clinical trials that are subject to FDAAA, responsible parties must register once,

not later than 21 days after enrolling the first subject. Updates to submitted information are required at least once a year, if there are changes to report, although changes in recruitment status and completion of a trial must be reported not later than 30 days after such events. Results information is to be submitted not later than 12 months after the completion date (as defined in the law), but can be delayed under certain circumstances. Other clinical studies also register once, at their inception, and are requested to update information annually, as necessary. An organization must establish a PRS account one time in order to register studies (and submit results) with ClinicalTrials.gov.

Description of Respondents: Respondents include sponsors or principal investigators of clinical studies. Those subject to FDAAA are referred to as "responsible parties," which are defined as sponsors of the clinical trial (as defined in 21 CFR 50.3) or designated principal investigators who meet requirements specified in the law. **Estimate of Burden:** The burden associated with this information collection is calculated in three parts: the burden associated with the one-time process of applying for a PRS account at ClinicalTrials.gov; the burden associated with registration; and the burden associated with the submission of results information, including adverse events. These information collections will occur at different times, but the registration and results information will be integrated into a single record for each clinical trial, which is entered through the PRS account. Based on data from 2011, we estimate that 5,500 new PRS account applications will be submitted annually. The time necessary to collect the required information and enter it into a new application form is estimated at 15 minutes. Using these figures, we estimate the total annual burden of submitting an application for a new PRS account to be 1,375 hours (5,500 applications per year times 0.25 hours per application). To estimate the annual reporting burden for registration, we examined the number of clinical studies registered annually with ClinicalTrials.gov and found an average of 17,000 registrations per year since the enactment of FDAAA. From this total, we estimate that approximately 5,000 studies would be applicable clinical trials of drugs (including biological products) and 500 would be applicable trials of devices subject to FDAAA. The remaining 11,500 studies would be registered voluntarily. We estimate the time to complete an initial registration