public docket files. The Federal Reserve 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.

FOR FURTHER INFORMATION CONTACT: Federal Reserve Board Clearance Officer—Michelle Shore—Division of Research and Statistics, Board of Governors of the Federal Reserve System, Washington, DC 20551 (202–452–3829)

OMB Desk Officer—Alexander T. Hunt—Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Final approval under OMB delegated authority of the extension for three years, without revision, of the following reports:

1. *Report Title:* Recordkeeping and Disclosure Requirements Associated with the Guidance on Response Programs for Unauthorized Access to Customer Information.

Agency Form Number: FR 4100.

OMB Control Number: 7100–0309.

Frequency: Develop customer notice, one-time; update and maintain customer notice, annually; Incident notification, event-generated.

Reporters: Financial institutions.

Annual Reporting Hours: 62,135.

Estimated Average Hours per Response: Develop customer notice, 24; Update and maintain customer notice, 8; Incident notification, 29.

Number of Respondents: Develop customer notice, 102; Update and maintain customer notice, 6,957; Incident notification, 139.

General Description of Report: This information collection is mandatory (15 U.S.C. 6801(b)). Since the Federal Reserve does not collect information associated with the FR 4100, any issue of confidentiality would not generally be an issue. However, confidentiality may arise if the Federal Reserve were to obtain a copy of a customer notice during the course of an examination or were to receive a copy of a Suspicious Activity Report (SAR; FR 2230; OMB No. 7100–0212). In such cases the information would be exempt from disclosure to the public under the Freedom of Information Act (5 U.S.C 552(b)(3), (4), and (8)). Also, a federal employee is prohibited by law from disclosing an SAR or the existence of an SAR (31 U.S.C. 5318(g)).

Abstract: Recent trends in customer information theft and the accompanying misuse of that information have led to the issuance of a supplemental interpretation of existing information technology-related security guidelines applicable to financial institutions. The supplemental guidelines are designed to facilitate timely and relevant notification of affected customers and the appropriate regulatory authority of the financial institutions. The guidelines provide specific direction regarding the nature and content of customer notice.

Current Actions: On March 6, 2008, the Federal Reserve published a notice in the **Federal Register** (73 FR 12176) requesting public comment for sixty days on the extension, without revision, of the ID-Theft Guidance. The comment period for this notice expired on May 5, 2008. The Federal Reserve did not receive any comments.

2. *Report Title:* The Recordkeeping and Disclosure Requirement in Connection with Regulation DD (Truth in Savings).

Agency Form Number: Reg DD.

OMB Control Number: 7100–0271.

Frequency: Account disclosures, 500; Change in terms notices, 1,130; Prematurity notices, 1,015; Disclosures on periodic statements, 12; and Advertising, 12.

Reporters: State member banks.

Annual Reporting Hours: 176,177.

Estimated Average Hours per Response: Account disclosures, 1.5 minutes; Change in terms notices, 1 minute; Prematurity notices, 1 minute; Disclosures on periodic statements, 8 hours; and Advertising, 30 minutes.

Number of Respondents: 1,172.

General Description of Report: This information collection is mandatory (12 U.S.C. 4308)). Since the Federal Reserve does not collect any information, no issue of confidentiality arises.

Abstract: The Truth in Savings Act and Regulation DD require depository institutions to disclose yields, fees, and other terms concerning deposit accounts to consumers at account opening, upon request, and when changes in terms occur. Depository institutions that provide periodic statements are required to include information about fees imposed, interest earned, and the annual percentage yield (APY) earned during those statement periods. The act and regulation mandate the methods by which institutions determine the account balance on which interest is calculated. They also contain rules about advertising deposit accounts.

Current Actions: On March 6, 2008, the Federal Reserve published a notice in the **Federal Register** (73 FR 1276) requesting public comment for sixty days on the extension, without revision, of the recordkeeping and disclosure requirements of Regulation DD. The comment period for this notice expired on May 5, 2008. The Federal Reserve did not receive any comments.

Board of Governors of the Federal Reserve System, May 9, 2008.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. E8–10780 Filed 5–14–08; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisition of Shares of Bank or Bank Holding Companies

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the office of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than May 30, 2008.

A. Federal Reserve Bank of Chicago (Burl Thornton, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690–1414:

1. Mark R. Peterson, Dakota Dunes, South Dakota, to acquire control of Liberty Financial Services, Inc., and thereby indirectly Liberty National Bank, both of Sioux City, Iowa.

Board of Governors of the Federal Reserve System, May 12, 2008.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E8–10858 Filed 5–14–08; 8:45 am]

BILLING CODE 6210–01–S

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 also will 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. Additional information on all bank holding companies may be obtained from the National Information Center website at *www.ffiec.gov/nic/*.

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 June 9, 2008.

A. Federal Reserve Bank of Chicago (Burl Thornton, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690–1414:

1. Premier Bancorp of Illinois, Inc., Farmer City, Illinois, to retain 20.8 percent of the voting shares of F M Bancorp, Inc., and thereby indirectly retain voting shares of Farmers–Merchants National Bank of Paxton, both of Paxton, Illinois.

B. Federal Reserve Bank of San Francisco (Kenneth Binning, Director, Regional and Community Bank Group) 101 Market Street, San Francisco, California 94105–1579:

1. Bank of Whitman Employee Stock Ownership Plan, Colfax Washington, to acquire 56 percent of the voting shares of Whitman Bancorporation Incorporated, Colfax, Washington, and thereby indirectly acquire voting shares of Bank of Whitman, Colfax, Washington.

Board of Governors of the Federal Reserve System, May 12, 2008.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc.E8–10859 Filed 5–14–08; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

In accordance with section 10(d) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2), announcement is made of a Health Care Policy and Research Special Emphasis Panel (SEP) meeting.

A Special Emphasis Panel is a group of experts in fields related to health care research who are invited by the Agency for Healthcare Research and Quality (AHRQ), and agree to be available, to conduct on an as needed basis, scientific reviews of applications for AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

Substantial segments of the upcoming SEP meeting listed below will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6). Grant applications for the Announcement of Availability of Funds for Grants regarding Ambulatory Safety and Quality Program: Improving Management of Individuals with Complex Healthcare Needs through Health IT (R18) applications are to be reviewed and discussed at this meeting. These discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure under the above-cited statutes.

SEP Meeting on: Ambulatory Safety and Quality Program: Improving Management of Individuals with Complex Healthcare Needs through Health IT (R18).

Date: June 18–20, 2008 (Open on June 18 from 5 p.m. to 5:15 p.m. and closed for the remainder of the meeting).

Place: Crowne Plaza, Conference Room TBD, 3 Research Blvd, Rockville, Maryland 20850.

Contact Person: Anyone wishing to obtain a roster of members, agenda or minutes of the nonconfidential portions of this meeting should contact Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, AHRQ, 540 Gaither Road, Room 2038, Rockville, Maryland 20850, Telephone (301) 427–1554.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: May 5, 2008.

Carolyn M. Clancy,

Director.

[FR Doc. E8–10565 Filed 5–14–08; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2007–E–0035 (formerly Docket No. 2007E–0133) and [Docket No. FDA–2007–E–0227 (formerly Docket No. 2007E–0148)]

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

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for TYZEKA and is publishing this notice of that determination as required by law. FDA has made the determination because of the submissions of applications to the Director of Patents and Trademarks, Department of Commerce, for the extension of patents which claim 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.regulations.gov*.

FOR FURTHER INFORMATION CONTACT: Beverly Friedman, Office of Regulatory Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg 51, rm. 6222, Silver Spring, MD 20993–0002, 301–796–3602.

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