President) 701 East Byrd Street, Richmond, Virginia 23261–4528:≤

1. Peden B. McLeod, Mary H. McLeod, John R. McLeod, all of Walterboro, South Carolina; Peden B. McLeod, Jr., Mt. Pleasant, South Carolina; Mary C. Benson, Columbia, South Carolina; and Rhoda L. Perry, Hendersonville, North Carolina; acting in concert to retain voting shares of Communitycorp, and thereby indirectly retain voting shares of Bank of Walterboro, both of Walterboro, South Carolina.

B. Federal Reserve Bank of St. Louis (Glenda Wilson, Community Affairs Officer) P.O. Box 442, St. Louis, Missouri 63166–2034:

1. First State Bancorp, Inc. Employee Stock Ownership Plan (Irvin G. Waller and Duane S. Michie as trustees), all of Caruthersville, Missouri; to acquire voting shares of First State Bancorp, Inc., and thereby indirectly acquire voting shares of First State Bank and Trust Company, both of Caruthersville, Missouri.

Board of Governors of the Federal Reserve System, March 11, 2010.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 2010–5686 Filed 3–15–10; 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 April 9, 2010.

A. Federal Reserve Bank of Philadelphia (Michael E. Collins, Senior Vice President) 100 North 6th Street, Philadelphia, Pennsylvania 19105–1521:

1. Tower Bancorp, Inc., Harrisburg, Pennsylvania; to merge with First Chester County Corporation, and thereby indirectly acquire First National Bank of Chester County, both of West Chester, Pennsylvania.

B. Federal Reserve Bank of St. Louis (Glenda Wilson, Community Affairs Officer) P.O. Box 442, St. Louis, Missouri 63166–2034:

1. Du Quoin State Bank ESOP, Du Quoin, Illinois; to become a bank holding company by retaining voting shares of Perry County Bancorp, Inc., and Du Quoin State Bank, both of Du Quoin, Illinois.

Board of Governors of the Federal Reserve System, March 11, 2010.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 2010–5687 Filed 3–15–10; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, e-mail *paperwork@hrsa.gov* or call the HRSA Reports Clearance Officer at (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: Ryan White Treatment and Modernization Act Part A Minority AIDS Initiative Report (the Part A MAI Report). (OMB No. 0915–0304): Extension

HRSA's HIV/AIDS Bureau (HAB) administers Part A of Title XXVI of the Public Health Service Act as amended by Congress in October 2009 (Ryan White HIV/AIDS Treatment Extension Act of 2009). Part A provides emergency relief for areas with substantial need for HIV/AIDS care and support services that are most severely affected by the HIV/AIDS epidemic, including eligible metropolitan areas (EMA) and Transitional Grant Areas (TGAs). As a component of Part A (previously Title I), the purpose of the Minority AIDS Initiative (MAI) Supplement is to improve access to high quality HIV care services and health outcomes for individuals in disproportionately impacted communities of color who are living with HIV disease, including African-Americans, Latinos, Native Americans, Asian Americans, Native Hawaiians and Pacific Islanders (Section 2693(b)(2)(A) of the Public Health Service (PHS) Act). Since the purpose of the Part A MAI is to expand access to medical, health, and social support services for disproportionately impacted racial/ethnic minority populations living with HIV/AIDS, who are not yet in care, it is important that HRSA is able to report on minorities served by the Part A MAI.

The Part A MAI Report is a data collection instrument in which grantees report on the number and characteristics of clients served and services provided. The *Part A MAI Report,* first approved for use in March 2006, is designed to collect performance data from Part A Grantees that will not change, and it has two parts: (1) a web-based data entry application that collects standardized quantitative and qualitative information, and (2) an accompanying narrative report. Grantees submit two *Part A MAI Reports* annually: *Part A MAI Plan* (*Plan*) and the *Part A MAI Year-End Annual Report* (*Annual Report*). The *Plan* and *Annual Report* components of the report are linked to minimize the reporting burden, and include drop-down menu responses, fields for reporting budget, expenditure and aggregated client level data, and open-ended responses for describing client or service-level outcomes. Together the *Plan* and *Annual Report* components collect information from grantees on MAI-funded services, expenditure patterns, the number and demographics of clients served, and client-level outcomes.

The MAI *Plan* Narrative that accompanies the *Plan* Web forms provides (1) an explanation of the data submitted in the *Plan* Web forms; (2) a summary of the *Plan,* including the plan and timeline for disbursing funds, monitoring service delivery, and implementing any service-related capacity development or technical assistance activities; and (3) the plan and timeline for documenting client-level outcome measures. In addition, if the EMA/TGA revised any planned services, allocation amounts or target communities after their grant application was submitted, the changes must be highlighted and explained. The accompanying MAI *Annual Report* Narrative describes (1) progress towards achieving specific goals and objectives identified in the Grantee's approved *MAI Plan* for that fiscal year and in linking MAI services/activities to Part A and other Ryan White HIV/AIDS Program services; (2) achievements in relation to client-level health outcomes; (3) summary of challenges or barriers at the provider or grantee levels, the strategies and/or action steps implemented to address them, and lessons learned; and, (4) discussion of MAI technical assistance needs identified by the EMA/TGA.

This information is needed to monitor and assess: (1) Changes in the type and amount of HIV/AIDS health care and related services being provided to each disproportionately impacted community of color; (2) the aggregate number of persons receiving HIV/AIDS services within each racial and ethnic community; and (3) the impact of Part A MAI-funded services in terms of client-level and service-level health outcomes. The information also is used to plan new technical assistance and capacity development activities, and inform the HRSA policy and program management functions. The data provided to HRSA does not contain individual or personally identifiable information.

The annual estimated response burden for grantees is as follows:

Form	Estimated number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Part A MAI Report	56	2	112	5	560

Note: Data collection system enhancements have resulted in a shortened response burden (from 6 to 5 total hours per response) for respondents since the previous OMB approval request.

E-mail comments to *paperwork@hrsa.gov* or mail the HRSA Reports Clearance Officer, Room 10–33, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Written comments should be received within 30 days of this notice.

Dated: March 5, 2010.

Sahira Rafiullah,

Director, Division of Policy and Information Coordination.

[FR Doc. 2010–5673 Filed 3–15–10; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0120]

Agency Information Collection Activities; Proposed Collection; Comment Request; Cosmetic Labeling Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection provisions in FDA's cosmetic labeling regulations.

DATES: Submit written or electronic comments on the collection of information by May 17, 2010.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Cosmetic Labeling Regulations—21 CFR Part 701 (OMB Control Number 0910–0599)—Extension

The Federal Food, Drug, and Cosmetic Act (the act) and the Fair Packaging and Labeling Act (the FPLA) require that cosmetic manufacturers, packers, and distributors disclose information about themselves or their products on the labels or labeling of their products.