FEDERAL MARITIME COMMISSION

Notice of Agreement(s) Filed

The Commission hereby gives notice of the filing of the following agreement(s) under the Shipping Act of 1984.

Interested parties can review or obtain copies of agreements at the Washington, DC offices of the Commission, 800 North Capitol Street, NW, Room 962. Interested parties may submit comments on an agreement to the Secretary, Federal Maritime Commission, Washington, DC 20573, within 10 days of the date of this notice appears in the **Federal Register**.

Agreement No.: 203–011560–001
Title: The TransAtlantic Bridge Agreement
Parties:
The COSCO/KL TransAtlantic Vessel Sharing Agreement (FMC Agreement No. 232–011561)
The KL/YM TransAtlantic Vessel Sharing Agreement (FMC Agreement No. 232–011562)
Synopsis: The proposed amendment would extend the term of the Agreement to October 31, 2000.

Agreement No.: 203–011561–001
Title: The COSCO/KL TransAtlantic Vessel Sharing Agreement
Parties:
China Ocean Shipping (Group) Company (''COSCO'')
Kawasaki Kisen Kaisha, Ltd. (''KL'')
Synopsis: The proposed amendment would extend the term of the Agreement to October 31, 2000.

Agreement No.: 232–011562–002
Title: The KL/YM TransAtlantic Vessel Sharing Agreement
Parties:
Yangming Transportation Corporation (''YM'') Company (''COSCO'')
Kawasaki Kisen Kaisha, Ltd. (''KL'')
Synopsis: The proposed amendment would extend the term of the Agreement to October 31, 2000.

Dated: October 7, 1998.

By Order of the Federal Maritime Commission.

Joseph C. Polking,

Secretary.

[FR Doc. 98–27411 Filed 10–13–98; 8:45 am]

BILLING CODE 6730–01–M

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of Banks 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 offices 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 October 27, 1998.

A. Federal Reserve Bank of St. Louis (Randall C. Sumner, Vice President) 411 Locust Street, St. Louis, Missouri 63102-2034:

1. First Perry Bancorp ESOP, Pinckneyville, Illinois; to acquire additional voting shares of First Perry Bancorp, Inc., Pinckneyville, Illinois, and thereby indirectly acquire First National Bank in Pinckneyville, Pinckneyville, Illinois.

Board of Governors of the Federal Reserve System, October 7, 1998.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–27505 Filed 10–13–98; 8:45 am]

BILLING CODE 6210–01–F

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 application 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. 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 November 6, 1998.

A. Federal Reserve Bank of Atlanta (Lois Berthaume, Vice President) 104 Marietta Street, N.W., Atlanta, Georgia 30303-2713:

1. First Community Bancorp, Inc., Pahokee, Florida; to become a bank holding company by acquiring 100 percent of the voting shares of First Community Bank of Palm Beach County, Pahokee, Florida.

B. Federal Reserve Bank of Kansas City (D. Michael Manies, Assistant Vice President) 925 Grand Avenue, Kansas City, Missouri 64198-0001:

1. Central Financial Corporation, Hutchinson, Kansas; to acquire 9.9 percent of the voting shares of Fort Worth National Bank, Fort Worth, Texas.

Board of Governors of the Federal Reserve System, October 7, 1998.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–27504 Filed 10–13–98; 8:45 am]

BILLING CODE 6210–01–F

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 11:00 a.m., Monday, October 19, 1998.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, N.W., Washington, D.C. 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any items carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board; 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications

scheduled for the meeting; or you may contact the Board's Web site at http://www.federalreserve.gov for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: October 9, 1998.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–27732 Filed 10–9–98; 3:44 pm]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0494]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Medical Device Registration and Listing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Submit written comments on the collection of information by November 13, 1998.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In compliance with section 3507 of the PRA (44 U.S.C. 3507), FDA has submitted the following proposed collection of information to OMB for review and clearance.

Medical Device Registration and Listing—21 CFR 807

Section 510 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360) requires that manufacturers and initial importers engaged in the manufacture, preparation, propagation, compounding, assembly, or processing of medical devices intended for human use and in commercial distribution register their establishments and list the devices they manufacture with FDA. This is accomplished by completing FDA Form 2891, ''Initial Registration of Device Establishment,'' and FDA Form 2892, ''Medical Device Listing.'' In addition, each year active, registered establishments must notify FDA of changes to the current registration and device listing for the establishment. Annual changes to current registration information are pre-printed on FDA Form 2891a and sent to registered establishments. The form must be sent back to FDA's Center for Devices and Radiological Health (CDRH), even if no changes have occurred. Changes to listing information are submitted on Form 2892. Refurbishers/reconditioners are not required to register or list; however, FDA will accept voluntary registration and listings from firms that wish to be registered with FDA.

In addition, under § 807.31 (21 CFR 807.31), each owner or operator is required to maintain a historical file containing the labeling and advertisements in use on the date of initial listing, and in use after October 10, 1978, but before the date of initial listing. The owner or operator must maintain in the historical file any labeling or advertisements in which a material change has been made anytime after initial listing, but may discard labeling and advertisements from the file 3 years after the date of the last shipment of a discontinued device by an owner or operator. Along with the recordkeeping requirements, the owner or operator must be prepared to submit to FDA upon specific request all labeling and advertising mentioned in the previous paragraph (§ 807.31(e)).

The information collected through these provisions is used by FDA to identify firms subject to the agency's regulations and is used to identify geographic distribution in order to effectively allocate its field resources for these inspections and to identify the class of the device which determines the inspection frequency. When complications occur with a particular device or component, manufacturers of similar or related devices can easily be identified.

The likely respondents to this information collection will be domestic establishments engaged in the manufacture, preparation, propagation, compounding, assembly, or processing of medical devices intended for human use and commercial distribution.

In the **Federal Register** of July 16, 1998 (63 FR 38409), the agency requested comments on the proposed collections of information. No significant comments were received.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	FDA Form	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
807.22(a)	Form 2891—Initial Establishment, Registration	1,462	1	1,462	.25	366
807.22(b)	Form 2892—Device Listing (initial and update)	5,640	1	5,640	.50	2,820
807.22(a)	Form 2891(a)—Registration Update	22,000	1	22,000	.25	5,500
807.31(e)		200	1	200	.50	100
Total						8,786

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.