Connection also will carry the meeting live via the Internet. To purchase these services call (703) 993–3100 or go to *http://www.capitolconnection.gmu.edu*.

Copies of materials adopted at this meeting can be purchased from the FCC's duplicating contractor, Best Copy and Printing, Inc. (202) 488–5300; Fax (202) 488–5563; TTY (202) 488–5562. These copies are available in paper format and alternative media, including large print/type; digital disk; and audio and video tape. Best Copy and Printing, Inc. may be reached by e-mail at *FCC@BCPIWEB.com*.

Federal Communications Commission.

Marlene H. Dortch,

Secretary.

[FR Doc. E8–29910 Filed 12–12–08; 4:15 pm]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

Open Commission Meeting Scheduled for Thursday, December 18, 2008, Canceled

December 12, 2008.

The Federal Communications Commission has canceled the Open Meeting on the items listed below, previously scheduled for Thursday, December 18, 2008, at 445 12th Street, SW., Washington, DC. The following items will remain on circulation for the Commissioners to vote:

Advanced Wireless Services (AWS)

• A Report and Order and Order on Reconsideration addressing service rules for fixed and mobile services, including advanced wireless services (AWS), in the 2155–2180 MHz band (AWS–3).

Program Carriage and Program Access

• A Report and Order modifying the program carriage rules and procedures and a Further Notice of Proposed Rulemaking seeking comment on the practices of programmers and broadcasters.

DTV Translators

• A Notice of Proposed Rulemaking proposing a new digital television translator service for analog loss areas.

DTV Consumer Education Notice of Apparent Liability

• An omnibus NAL against various companies for apparent violations of the Commission's DTV consumer education requirements.

E911 Location Accuracy Requirements

• A Second Report and Order addressing the geographic area over which wireless carriers must meet the Enhanced 911 (E911) location accuracy requirements.

Regulatory Framework for SDARS and WCS in the 2305–2360 MHz Band

• A Report and Order and Second Report and Order and Order addressing the a regulatory framework for the coexistence of licensees in the Satellite Digital Audio Radio Service (SDARS) and the Wireless Communications Service (WCS) in the 2305–2360 MHz frequency band.

Wireless Radio Services (WRS) Renewals

• A Notice of Proposed Rulemaking and Order addressing Amendment of Parts 1, 22, 24, 27, 74, 80, 90, 95, and 101 To Establish License Renewal and Discontinuance of Operation Policies and Procedures for Certain Wireless Radio Services; Imposition of a Freeze on the Filing of Competing Renewal Applications for Certain Wireless Radio Services and the Processing of Already-Filed Competing Renewal Applications.

Additional information concerning this meeting may be obtained from Audrey Spivack or David Fiske, Office of Media Relations, (202) 418–0500; TTY 1–888–835–5322.

Federal Communications Commission.

Marlene H. Dortch,

Secretary.

[FR Doc. E8–30100 Filed 12–15–08; 4:15 pm]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

Notice of Agreement Filed

The Commission hereby gives notice of the filing of the following agreement under the Shipping Act of 1984. Interested parties may submit comments on agreements to the Secretary, Federal Maritime Commission, Washington, DC 20573, within ten days of the date this notice appears in the **Federal Register**. Copies of agreements are available through the Commission's Web site (*http://www.fmc.gov*) or contacting the Office of Agreements at (202) 523–5793 or *tradeanalysis@fmc.gov*.

Agreement No.: 012059.

Title: CSAV/NYK Venezuela Space Charter Agreement.

Parties: Compania Sud Americana De Vapores S.A. and Nippon Yusen Kaisha.

Filing Party: Michael B. Holt, Vice President and General Counsel; NYK Line; 300 Lighting Way, 5th Floor; Secaucus, NJ 07094.

Synopsis: The agreement authorizes CSAV to charter space to NYK in the trade from Newark, NJ, Baltimore, MD, and Miami, FL, to ports in Venezuela.

By Order of the Federal Maritime Commission.

Dated: December 12, 2008.

Tanga S. Fitzgibbon,

Alternate Federal Register Liaison Officer.

[FR Doc. E8–29945 Filed 12–16–08; 8:45 am]

BILLING CODE 6730–01–P

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 January 12, 2009.

A. Federal Reserve Bank of New York (Anne MacEwen, Bank Applications Officer) 33 Liberty Street, New York, New York 10045-0001:

1. Morgan Stanley, New York, New York; to acquire additional common

shares up to 9.9 percent of the voting shares of Chinatrust Financial Holding Company, Ltd., Taipei, Taiwan, and thereby acquire Chinatrust Bank (U.S.A.), Torrance, California.

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

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E8–29914 Filed 12–16–08; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Notice of Proposals to Engage in Permissible Nonbanking Activities or to Acquire Companies that are Engaged in Permissible Nonbanking Activities

The companies listed in this notice have given notice under section 4 of the Bank Holding Company Act (12 U.S.C. 1843) (BHC Act) and Regulation Y (12 CFR Part 225) to engage *de novo*, or to acquire or control voting securities or assets of a company, including the companies listed below, that engages either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.28 of Regulation Y (12 CFR 225.28) or that the Board has determined by Order to be closely related to banking and permissible for bank holding companies. Unless otherwise noted, these activities will be conducted throughout the United States.

Each notice is available for inspection at the Federal Reserve Bank indicated. The notice also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the question whether the proposal complies with the standards of section 4 of the BHC Act. 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 the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than January 12, 2008.

A. Federal Reserve Bank of Richmond (A. Linwood Gill, III, Vice President) 701 East Byrd Street, Richmond, Virginia 23261-4528:

1. Capital One Financial Corporation, McLean, Virginia; to acquire Chevy Chase Bank, Federal Savings Bank, McLean, Virginia, and thereby engage in operation a savings and loan association pursuant to section 225.28(b)(4)(ii) of Regulation Y.

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

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E8–29913 Filed 12–16–08; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0439]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Blood Establishment Registration and Product Listing, Form FDA 2830

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

DATES: Fax written comments on the collection of information by January 16, 2009.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0052. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794.

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

Blood Establishment Registration and Product Listing, Form FDA 2830—(OMB Control Number 0910–0052)—Extension

Under section 510 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360), any person owning or operating an establishment that manufactures, prepares, propagates, compounds, or processes a drug or device must register with the Secretary of Health and Human Services, on or before December 31 of each year, his or her name, place of business, and all such establishments, and must submit, among other information, a listing of all drug or device products manufactured, prepared, propagated, compounded, or processed by him or her for commercial distribution. In part 607 (21 CFR part 607), FDA has issued regulations implementing these requirements for manufacturers of human blood and blood products.

Section 607.20(a), in brief, requires owners or operators of certain establishments that engage in the manufacture of blood products to register and to submit a list of every blood product in commercial distribution. Section 607.21, in brief, requires the owners or operators of establishments entering into the manufacturing of blood products to register within 5 days after beginning such operation and to submit a list of every blood product in commercial distribution at the time. If the owner or operator of the establishment has not previously entered into such operation for which a license is required, registration must follow within 5 days after the submission of a biologics license application. In addition, establishments are required to register annually between November 15 and December 31 and update their blood product listing every June and December of each year. Section 607.22 requires the use of Form FDA 2830, Blood Establishment Registration and Product Listing, for initial registration, for annual registration, and for blood product listing. Section 607.25 indicates the information required for establishment registration and blood product listing. Section 607.26, in brief, requires certain changes to be submitted on FDA Form 2830 as amendments to the establishment registration within 5 days of such changes. Section 607.30(a), in brief, indicates the information required for owners or operators of establishments to update their blood product listing information every June and December, or at the discretion of the registrant at the time the change occurs. Section 607.31 requires that additional blood product listing information be provided upon FDA request. Section 607.40, in brief, requires certain foreign blood product establishments to register and submit the blood product listing information, and to provide the name and address of the establishment and the name of the individual responsible for submitting blood product listing