(d)(1) Time brokerage agreements (also known as local marketing agreements): Time brokerage agreements involving radio stations where the licensee (including all parties under common ownership) is the brokering entity, the brokering and brokered stations are both in the same market as defined in the local radio multiple ownership rule contained in §73.3555(a), and more than 15 percent of the time of the brokered station, on a weekly basis is brokered by that licensee; time brokerage agreements involving television stations where the licensee (including all parties under common control) is the brokering entity, the brokering and brokered stations are both licensed to the same market as defined in the local television multiple ownership rule contained in §73.3555(b), and more than 15 percent of the time of the brokered station, on a weekly basis, is brokered by that licensee; time brokerage agreements involving radio or television stations that would be attributable to the licensee under §73.3555 Note 2, paragraph (i). Confidential or proprietary information may be redacted where appropriate but such information shall be made available for inspection upon request by the FCC.

(2) Joint sales agreements: Joint sales agreements involving radio stations where the licensee (including all parties under common control) is the brokering entity, the brokering and brokered stations are both in the same market as defined in the local radio multiple ownership rule contained in §73.3555(a), and more than 15 percent of the advertising time of the brokered station on a weekly basis is brokered by that licensee. Confidential or proprietary information may be redacted where appropriate but such information shall be made available for inspection upon request by the FCC.

(e) The following contracts, agreements or understandings need not be filed but shall be kept at the station and made available for inspection upon request by the FCC; subchannel leasing agreements for Subsidiary Communications Authorization operation; franchise/leasing agreements for operation of telecommunications services on the television vertical blanking interval and in the visual signal; time sales contracts with the same sponsor for 4 or more hours per day, except where the length of the events (such as athletic contests, musical programs and special events) broadcast pursuant to the contract is not under control of the station; and contracts with chief operators.

Federal Communications Commission.

Marlene H. Dortch,

Secretary,

Office of the Secretary,

Office of Managing Director.

[FR Doc. 2010–22250 Filed 9–7–10; 8:45 am]

BILLING CODE 6712–01–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 October 4, 2010.

A. Federal Reserve Bank of Chicago, (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. Zaring Group Holdings LLC., Riverwoods, Illinois, to become a bank holding company by acquiring 75.1 percent of the voting shares of First Suburban Bancorp Corporation, Maywood, Illinois, and thereby indirectly acquire First Suburban National Bank, Maywood, Illinois.

2. Hometown Community Bancorp, Inc., and Hometown Community Bancorp, Inc. Employee Stock Ownership Plan and Trust, both located in Morton, Illinois, to merge with CSBC Financial Corporation, Cropsey, Illinois, and thereby indirectly acquire Citizens State Bank of Cropsey, Cropsey, Illinois.

Board of Governors of the Federal Reserve System, September 2, 2010.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2010–22310 Filed 9–7–10; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Chronic Fatigue Syndrome Advisory Committee

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services is hereby giving notice that the Chronic Fatigue Syndrome Advisory Committee (CFSAC) will hold a meeting. The meeting will be open to the public.

DATES: The meeting will be held on Wednesday, October 13 from 9 a.m. until 5 p.m. and Thursday, October 14, 2010 from 9 a.m. until 4 p.m. CFSAC Subcommittees will hold scientific review sessions on Tuesday, October 12 from 8:30 a.m. until 5 p.m.

ADDRESSES: Department of Health and Human Services; Room 800, Hubert H. Humphrey Building; 200 Independence Avenue, SW., Washington, DC 20201. For a map and directions to the Hubert H. Humphrey building, please visit *http://www.hhs.gov/about/hhhmap.html.*

FOR FURTHER INFORMATION CONTACT: Wanda K. Jones, DrPH; Executive Secretary, Chronic Fatigue Syndrome Advisory Committee, Department of Health and Human Services; 200 Independence Avenue, SW., Hubert Humphrey Building, Room 712E; Washington, DC 20201. Please direct all inquiries to *cfsac@hhs.gov.*

SUPPLEMENTARY INFORMATION: CFSAC was established on September 5, 2002. The Committee was established to advise, consult with, and make recommendations to the Secretary, through the Assistant Secretary for Health, on a broad range of topics including (1) The current state of the knowledge and research about the

epidemiology and risk factors relating to chronic fatigue syndrome, and identifying potential opportunities in these areas; (2) current and proposed diagnosis and treatment methods for chronic fatigue syndrome; and (3) development and implementation of programs to inform the public, health care professionals, and the biomedical, academic, and research communities about advances in chronic fatigue syndrome.

The agenda for this meeting is being developed. The agenda will be posted on the CFSAC Web site, *http://www.hhs.gov/advcomcfs,* when it is finalized. The meeting will be broadcast over the Internet as a real-time streaming video. It also will be recorded and archived on the CFSAC Web site for on demand viewing.

CFSAC Subcommittees will convene scientific review sessions on Tuesday, October 12. The purpose of these sessions is to update the latest developments in etiology, natural history, clinical trials, and related areas for chronic fatigue syndrome. The public is welcome to attend these sessions, which are not a formal part of the Advisory Committee meeting. These sessions will be broadcast over the Internet as a real-time streaming video. It also will be recorded and archived on the CFSAC Web site for on demand viewing. An agenda will be posted on the CFSAC Web site when it becomes available.

Public attendance at the meeting is limited to space available. Individuals must provide a government-issued photo ID for entry into the building where the meeting is scheduled to be held. Those attending the meeting will need to sign-in prior to entering the meeting room. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person at *cfsac@hhs.gov* in advance.

Members of the public will have the opportunity to provide comment at the October 13–14 meeting if pre-registered. Individuals who wish to address the Committee during the public comment session must pre-register by Friday, September 17, 2010, via e-mail at *cfsac@hhs.gov.* Time slots for public comment will be available on a first-come, first-served basis. Public comment will be limited to five minutes per speaker; no exceptions will be made. Individuals registering for public comment should submit a copy of their testimony in advance to *cfsac@hhs.gov,* prior to the close of business on Friday, September 17, 2010.

Members of the public who wish to have printed material distributed to CFSAC members for review should submit one copy of the material to the Executive Secretary, at *cfsac@hhs.gov,* prior to close of business on September 17, 2010. Submissions are limited to five typewritten pages. Any written testimony submitted after this date will be available for inspection on-site and will be posted to the Web site after the meeting.

If you do not submit your written testimony prior to the close of business Friday, September 17, 2010, you may bring a copy of your written testimony to the meeting and present it to the CFSAC Executive Secretary. Your testimony will be included in a notebook that will be available for viewing by the public on a table at the back of the meeting room.

Please ensure that written testimony does not include any personal information including your personal mailing address and that it includes only your name, if you wish to be identified. If you wish to remain anonymous, please notify the CFSAC Executive Secretary upon submission of the materials to *cfsac@hhs.gov.*

Dated: August 31, 2010.

Wanda K. Jones,

Executive Secretary, Chronic Fatigue Syndrome Advisory Committee.

[FR Doc. 2010–22393 Filed 9–7–10; 8:45 am]

BILLING CODE 4150–42–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–D–0285]

Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document for Certain Percutaneous Transluminal Coronary Angioplasty (PTCA) Catheters; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of the guidance entitled "Class II Special Controls Guidance Document for Certain Percutaneous Transluminal Coronary Angioplasty (PTCA) Catheters." The guidance was developed as a special control to support the reclassification of PTCA catheters, other than cutting/scoring PTCA catheters, from class III (premarket approval) into class II (special controls). This guidance describes a means by which PTCA catheters, other than cutting/scoring PTCA catheters, may comply with the requirement of special controls for class II devices. Elsewhere in this issue of the **Federal Register**, FDA is publishing a final rule that codifies the reclassification of this device type from class III (premarket approval) into class II (special controls).

DATES: Submit written or electronic comments on this guidance at any time. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Submit written requests for single copies of the guidance document entitled "Class II Special Controls Guidance Document for Certain Percutaneous Transluminal Coronary Angioplasty (PTCA) Catheters" to the Division of Small Manufacturers, International, and Consumer Assistance (HFZ–220), Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 4613, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your request, or fax your request to 301–847–8149. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance.

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

FOR FURTHER INFORMATION CONTACT: Kathryn O'Callaghan, Center for Devices and Radiological Health (HFZ–450), Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301–796–6349.

SUPPLEMENTARY INFORMATION:

I. Background

This guidance document was developed as a special control guidance to support the reclassification of PTCA catheters, other than cutting/scoring PTCA catheters, into class II (special controls). The device is intended for balloon dilatation of a hemodynamically significant coronary artery or bypass graft stenosis in patients evidencing coronary ischemia for the purpose of improving myocardial perfusion, treatment of acute myocardial infarction, treatment of in-stent restenosis and/or post-deployment stent expansion. Cutting/scoring PTCA catheters (Product Code: NWX) remain in class III and are subject to premarket