

respondents to the FR 2910a would reduce reporting burden by 3,690 hours.

Board of Governors of the Federal Reserve System, May 15, 2000.

Jennifer J. Johnson,
Secretary of the Board.

[FR Doc. 00-12589 Filed 5-18-00; 8:45 am]

BILLING CODE 6210-01-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 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 (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 12, 2000.

A. Federal Reserve Bank of Chicago (Phillip Jackson, Applications Officer) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. *Antioch Holding Company*, Antioch, Illinois; to acquire 24.99 percent of the voting shares of Lakes Region Bancorp, Inc, Third Lake, Illinois, and thereby indirectly acquire Anchor Bank, Third Lake, Illinois.

B. Federal Reserve Bank of St. Louis (Randall C. Sumner, Vice President) 411

Locust Street, St. Louis, Missouri 63166-2034;

1. *Valley Capital Corporation*, Greenwood, Mississippi; to merge with State Capital Corporation, Brookhaven, Mississippi, and thereby indirectly acquire State Bank and Trust Company, Brookhaven, Mississippi.

Board of Governors of the Federal Reserve System, May 15, 2000.

Robert deV. Frierson,
Associate Secretary of the Board.

[FR Doc. 00-12588 Filed 5-18-00; 8:45 am]

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BOARD OF GOVERNORS OF THE FEDERAL RESERVE SYSTEM

Sunshine Act Meeting; Notice

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

TIME AND DATE: 10 a.m., Wednesday, May 24, 2000.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, NW., Washington, DC 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 matters 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: May 17, 2000.

Robert deV. Frierson,
Associate Secretary of the Board.

[FR Doc. 00-12744 Filed 5-17-00; 10:54 am]

BILLING CODE 6210-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00D-1278]

Draft Guidance for Industry on Female Sexual Dysfunction: Clinical Development of Drug Products for Treatment; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled "Female Sexual Dysfunction: Clinical Development of Drug Products for Treatment." The draft guidance is intended to provide recommendations to industry on the development of drug products for the treatment of female sexual dysfunction (FSD).

DATES: Submit written comments on the draft guidance by July 18, 2000. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Copies of this draft guidance for industry can be obtained on the Internet at <http://www.fda.gov/cder/guidance/index.htm>. Submit written requests for single copies of the draft guidance to the Drug Information Branch (HFD-210), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857. Send one self-addressed adhesive label to assist that office in processing your requests. Submit written comments on the draft guidance to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Lana L. Pauls, Center for Drug Evaluation and Research (HFD-580), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-4260.

SUPPLEMENTARY INFORMATION: FDA is announcing the availability of a draft guidance for industry entitled "Female Sexual Dysfunction: Clinical Development of Drug Products for Treatment." The draft guidance provides recommendations for sponsors designing clinical trials in support of new drug applications for the treatment of FSD. It includes recommendations on the appropriate definition of the patient population to be studied, inclusion and exclusion criteria, the use of scales and questionnaires to assess FSD, and

primary endpoints for trials of drug products.

This Level 1 draft guidance is being issued consistent with FDA's good guidance practices (62 FR 8961, February 27, 1997). The draft guidance represents the agency's current thinking on the development of drugs for the treatment of FSD. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirement of the applicable statute, regulations, or both.

Interested persons may submit to the Dockets Management Branch (address above) written comments on the draft guidance. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. The draft guidance and received comments are available for public examination in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Dated: May 12, 2000.

Margaret M. Dotzel,

Acting Associate Commissioner for Policy.
[FR Doc. 00-12594 Filed 5-18-00; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA-3034-N]

Medicare Program; Meeting of the Executive Committee of the Medicare Coverage Advisory Committee—June 6, 2000

AGENCY: Health Care Financing Administration (HCFA), HHS.

ACTION: Notice of meeting.

SUMMARY: This notice announces a public meeting of the Executive Committee (the Committee) of the Medicare Coverage Advisory Committee (MCAC). The Committee will hear reports from the Medical and Surgical Procedures Panel meeting of April 12 and 13, 2000, during which biofeedback and pelvic floor electrical stimulation in the treatment of urinary incontinence were deliberated. The Committee will also discuss presentations from interested persons regarding procedural aspects of future public meetings of the medical specialty panels of the MCAC. Notice of this meeting is given under the Federal Advisory Committee Act (5

U.S.C. App. 2, section 10(a)(1) and (a)(2)).

DATES:

The Meeting: June 6, 2000, from 8 a.m. until 3 p.m., E.D.T.

Deadline for Presentations and Comments: May 17, 2000, 5 p.m., E.D.T.

Special Accommodations: Persons attending the meeting who are hearing impaired and require sign language interpretation, or have a condition that requires other special assistance or accommodations, are asked to notify the Executive Secretary by May 25, 2000.

ADDRESSES:

The Meeting: The meeting will be held at the Baltimore Convention Center, One West Pratt Street, Baltimore, Maryland 21201.

Presentations and Comments: Submit formal presentations and written comments to Constance A. Conrad, Executive Secretary; Office of Clinical Standards and Quality; Health Care Financing Administration; 7500 Security Boulevard; Mail Stop S3-02-01; Baltimore, MD 21244.

Website: You may access up-to-date information on this meeting at www.hcfa.gov/quality/8b.htm.

Hotline: You may access up-to-date information on this meeting on the HCFA Advisory Committee Information Hotline, 1-877-449-5659 (toll free) or in the Baltimore area (410) 786-9379.

FOR FURTHER INFORMATION CONTACT:

Constance A. Conrad, Executive Secretary, 410-786-4631.

SUPPLEMENTARY INFORMATION: On August 13, 1999, we published a notice (64 FR 44231) to describe the MCAC, which provides advice and recommendations to us about clinical issues.

Current Panel Members

Harold C. Sox, MD (Chairperson); Thomas V. Holohan; Leslie P. Francis; John H. Ferguson; Robert L. Murray; Alan M. Garber; Michael D. Maves; Frank J. Papatheofanis; Ronald M. Davis; Daisy Alford-Smith; Joe W. Johnson; Robert H. Brook; Linda A. Bergthold; Randel E. Richner.

Meeting Topic

The Committee will hear reports from the Medical and Surgical Procedures Panel meeting of April 12 and 13, 2000, during which biofeedback and pelvic floor electrical stimulation in the treatment of urinary incontinence. The Committee will also hear and discuss presentations from interested persons regarding procedural aspects of future public meetings of the medical specialty panels of the MCAC.

Procedure and Agenda

This meeting is open to the public. The Committee will hear oral presentations from the public for approximately 1 hour. The Committee may limit the number and duration of oral presentations to the time available. If you wish to make formal presentations you must notify the For Further Information Contact, and submit the following by the Deadline for Presentations and Comments date listed in the **DATES** section of this notice: a brief statement of the general nature of the evidence or arguments you wish to present, the names and addresses of proposed participants, and an estimate of the time required to make the presentation. We will request that you declare at the meeting whether you have any financial involvement with manufacturers of any items or services being discussed (or with their competitors).

After the public presentation, we will make a presentation to the Committee. After our presentation, the Committee will deliberate openly. Interested persons may observe the deliberations, but the Committee will not hear further comments during this time except at the request of the chairperson. Prior to the review of the Medical Specialty Panel recommendation, the Committee will allow approximately a 30-minute open public session for any attendee to address issues specific to the topic. After the open session, the members will vote and the Committee will make its recommendation.

Authority: 5 U.S.C. App. 2, section 10(a)(1) and (a)(2).

(Catalog of Federal Domestic Assistance Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: May 11, 2000.

Jeffrey L. Kang,

Director Office of Clinical Standards and Quality, Health Care Financing Administration.

[FR Doc. 00-12686 Filed 5-18-00; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA-1136-N]

Medicare Program; June 5, 2000, Meeting of the Practicing Physicians Advisory Council

AGENCY: Health Care Financing Administration (HCFA), HHS.