SUMMARY: This notice sets forth the schedule and summary agenda for a meeting of the President's Council of Advisors on Science and Technology (PCAST), and describes the functions of the Council. Notice of this meeting is required under the Federal Advisory Committee Act (FACA).

Dates and Place: September 12, 2006, Washington, DC. The meeting will be held in the Continental Ballroom of the George Washington University Marvin Center Building, 800 21st St. NW., Washington DC 20052.

Type of Meeting: Open. Further details on the meeting agenda will be posted on the PCAST Web site at: *http://www.ostp.gov/PCAST/pcast.html.*

Proposed Schedule and Agenda: The President's Council of Advisors on Science and Technology is scheduled to meet in open session on Tuesday September 12, 2006, at approximately 9 a.m. The PCAST is tentatively scheduled to discuss its report and recommendations related to energy technology. The co-chairs of the PCAST subcommittee on networking and information technology are tentatively scheduled to provide an update on subcommittee activities and lead a discussion on the PCAST review of the Federal Networking and Information Technology Research and Development (NITRD) Program. A presentation on ethical and societal issues related to emerging technology capabilities is also tentatively scheduled to occur. This session will end at approximately 5 p.m. Additional information and the final agenda will be posted at the PCAST Web site at: *http://www.ostp.gov/PCAST/pcast.html.*

Public Comments: There will be time allocated for the public to speak on the above agenda items. This public comment time is designed for substantive commentary on PCAST's work topics, not for business marketing purposes. Please submit a request for the opportunity to make a public comment five (5) days in advance of the meeting. The time for public comments will be limited to no more than 5 minutes per person. Written comments are also welcome at any time following the meeting. Please notify Celia Merzbacher, PCAST Executive Director, at (202) 456–7116, or fax your request/comments to (202) 456–6021.

FOR FURTHER INFORMATION CONTACT: For information regarding time, place and agenda, please call Celia Merzbacher at (202) 456–7116, prior to 3 p.m. on Friday, September 8, 2006. Information will also be available at the PCAST Web site at: *http://www.ostp.gov/PCAST/pcast.html.* Please note that public seating for this meeting is limited and is available on a first-come, first-served basis.

SUPPLEMENTARY INFORMATION: The President's Council of Advisors on Science and Technology was established by Executive Order 13226, on September 30, 2001. The purpose of PCAST is to advise the President on matters of science and technology policy, and to assist the President's National Science and Technology Council in securing private sector participation in its activities. The Council members are distinguished individuals appointed by the President from non-Federal sectors. The PCAST is co-chaired by Dr. John H. Marburger, III, the Director of the Office of Science and Technology Policy, and by E. Floyd Kvamme, a Partner at Kleiner Perkins Caufield & Byers.

Celia Merzbacher,

PCAST Executive Director, Office of Science and Technology Policy.

[FR Doc. E6–14451 Filed 8–28–06; 8:45 am]

BILLING CODE 3170–W4–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 Web site at *http://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 September 25, 2006.

A. Federal Reserve Bank of Atlanta (Andre Anderson, Vice President) 1000 Peachtree Street, N.E., Atlanta, Georgia 30309:

1. Southcrest Financial Group, Inc., Fayetteville, Georgia; to merge with Maplesville Bancorp, and thereby indirectly acquire Peachtree Bank, both of Maplesville, Alabama.

B. Federal Reserve Bank of Minneapolis (Jacqueline G. King, Community Affairs Officer) 90 Hennepin Avenue, Minneapolis, Minnesota 55480-0291:

2. Baldwin Bancshares, Inc., Baldwin, Wisconsin; to acquire 100 percent of the voting shares of Gavic Services, Inc., Spring Valley, Wisconsin, and thereby indirectly acquire voting shares of The Bank of Spring Valley, Spring Valley, Wisconsin.

Board of Governors of the Federal Reserve System, August 24, 2006.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E6–14310 Filed 8–28–06; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

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

TIME AND DATE: 11:30 a.m., Tuesday, September 5, 2006.

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 items carried forward from a previously announced meeting.

FOR FURTHER INFORMATION CONTACT: Michelle Smith, Director, or Dave Skidmore, Assistant to the Board, Office of Board Members at 202–452–2955.

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.

Board of Governors of the Federal Reserve System, August 25, 2006.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 06–7253 Filed 8–25–06; 1:17 pm]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0203]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; User Fee Cover Sheet; Form FDA 3397

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 September 28, 2006.

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.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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.

User Fee Cover Sheet; Form FDA 3397—(OMB Control Number 0910–0297)—Extension

Under sections 735 and 736 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379g and 379h), the Prescription Drug User Fee Act of 1992 (PDUFA) (Public Law 102–571), as amended by the Food and Drug Administration Modernization Act of 1997 (Public Law 105–115), and the Public Health Security and Bioterrorism Preparedness and Response Act of 2002, which includes the Prescription Drug User Fee Amendments of 2002 (Public Law 107–188), FDA has the authority to assess and collect user fees for certain drug and biologics license applications and supplements. Under this authority, pharmaceutical companies pay a fee for certain new human drug applications, biologics license applications, or supplements submitted to FDA for review. Because the submission of user fees concurrently with applications and supplements is required, review of an application by FDA cannot begin until the fee is submitted. Form FDA 3397, the user fee cover sheet, is designed to provide the minimum necessary information to determine whether a fee is required for review of an application, to determine the amount of the fee required, and to account for and track user fees. The form provides a cross-reference of the fee submitted for an application with the actual application by using a unique number tracking system. The information collected is used by FDA's Center for Drug Evaluation and Research (CDER) and Center for Biologics Evaluation and Research (CBER) to initiate the administrative screening of new drug applications, biologics license applications, and supplemental applications.

Respondents to this collection of information are new drug and biologics manufacturers. Based on FDA's database system for fiscal year (FY) 2005, there are an estimated 243 manufacturers of products subject to PDUFA. However, not all manufacturers will have any submissions, and some may have multiple submissions in a given year. The total number of annual responses is based on the number of submissions received by FDA in FY 2005. CDER estimates 3,085 annual responses that include the following submissions: 101 new drug applications; 3 biologics license applications; 1,915 manufacturing supplements; 921 labeling supplements; and 145 efficacy supplements. CBER estimates 676 annual responses that include the following submissions: 6 biologics license applications, 614 manufacturing supplements, 46 labeling supplements, and 10 efficacy supplements. Based on previous estimates, the rate of submissions is not expected to change significantly in the next few years. The estimated hours per response are based on past FDA experience with the various submissions, and range from 5 to 30 minutes. The hours per response are based on the average of these estimates.

In the **Federal Register** of May 25, 2006 (71 FR 30144), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Form	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Respondent	Total Hours
FDA 3397	243	15.48	3,761	0.30	1,128

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