indicated. Once the application has been accepted for processing, it will also 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, including whether the acquisition of the nonbanking company can ''reasonably be expected to produce benefits to the public, such as greater convenience, increased competition, or gains in efficiency, that outweigh possible adverse effects, such as undue concentration of resources, decreased or unfair competition, conflicts of interests, or unsound banking practices'' (12 U.S.C. 1843). Any request for a hearing must be accompanied by a statement of the reasons a written presentation would not suffice in lieu of a hearing, identifying specifically any questions of fact that are in dispute, summarizing the evidence that would be presented at a hearing, and indicating how the party commenting would be aggrieved by approval of the proposal. 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 22, 1996.

A. Federal Reserve Bank of Atlanta (Zane R. Kelley, Vice President) 104 Marietta Street, N.W., Atlanta, Georgia 30303:

1. 1st Floyd Bankshares, Inc., Rome, Georgia; to become a bank holding company by acquiring 100 percent of the voting shares of 1st Floyd Bank, Rome, Georgia.

B. Federal Reserve Bank of Dallas (Genie D. Short, Vice President) 2200 North Pearl Street, Dallas, Texas 75201-2272:

1. MainBancorp, Inc., Austin, Texas, and MainCorp Intermediate Holding Company, Inc., Wilmington, Delaware; to become bank holding companies by acquiring 100 percent of the voting shares of ROSB Bancorp, Inc., Red Oak, Texas, and thereby indirectly acquire MainBank, Red Oak, Texas.

Board of Governors of the Federal Reserve System, October 23, 1996.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 96–27670 Filed 10–28–96; 8:45 am]

BILLING CODE 6210–01–F

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 that engages either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.25 of Regulation Y (12 CFR 225.25) 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. Once the notice has been accepted for processing, it will also 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, including whether consummation of the proposal can ''reasonably be expected to produce benefits to the public, such as greater convenience, increased competition, or gains in efficiency, that outweigh possible adverse effects, such as undue concentration of resources, decreased or unfair competition, conflicts of interests, or unsound banking practices'' (12 U.S.C. 1843). Any request for a hearing on this question must be accompanied by a statement of the reasons a written presentation would not suffice in lieu of a hearing, identifying specifically any questions of fact that are in dispute, summarizing the evidence that would be presented at a hearing, and indicating how the party commenting would be aggrieved by approval of the proposal.

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 November 12, 1996.

A. Federal Reserve Bank of Richmond (Lloyd W. Bostian, Jr., Senior Vice President) 701 East Byrd Street, Richmond, Virginia 23261:

1. MainStreet BankGroup Incorporated, Martinsville, Virginia; to engage *de novo* through its subsidiary, MainStreet Trust Company, NA, Martinsville, Virginia, in trust activities, pursuant to § 225.25(b)(3) of the Board's Regulation Y.

Board of Governors of the Federal Reserve System, October 23, 1996.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 96–27671 Filed 10–28–96; 8:45 am]

BILLING CODE 6210–01–F

Sunshine Meeting Notice

TIME AND DATE: 11:00 a.m., Monday, November 4, 1996.

PLACE: Marriner S. Eccles Federal Reserve Board Building, C Street entrance between 20th and 21st 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: Mr. Joseph R. Coyne, Assistant to the Board; (202) 452–3204. You may call (202) 452–3207, beginning at approximately 5 p.m. two business days before this meeting, for a recorded announcement of bank and bank holding company applications scheduled for the meeting.

Dated: October 25, 1996.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 96–27886 Filed 10–25–96; 3:23 pm]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 91N–0404]

Agency Information Collection; Submission for OMB Review; Comment Request

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.

DATES: Submit written comments on the collections of information by November 29, 1996.

ADDRESSES: Submit written comments on the collections of information to the

Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC, Attn: Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: In the Federal Register of June 26, 1996 (61 FR 33232), FDA issued a final rule implementing the provisions of the Safe Medical Devices Act (the SMDA) regarding Humanitarian Use Devices (HUD's). The final rule contained information collection requirements subject to the Paperwork Reduction Act of 1995 (the PRA) (44 U.S.C. 3507). In compliance with section 3507 of the PRA, FDA has submitted the following proposed collection of information to OMB for review and clearance:

Title: Medical Devices; Humanitarian Use Devices.

Description: This regulation implements the provision of the SMDA regarding HUD's. A HUD is exempt from the effectiveness requirements of sections 514 and 515 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360d and 360e). In order to implement this exemption, FDA is amending the premarket approval regulations in 21 CFR part 814 by creating new subpart H. This final regulation prescribes the procedures for submitting Humanitarian Device Exemption (HDE) applications, amendments and supplements; procedures for obtaining an extension of the exemption; and the criteria for FDA review and approval of HDE's. This final rule will create a needed incentive for the development of devices for use in the treatment or diagnosis of diseases or conditions affecting a small number of individuals.

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
814.102	20	1	20	40	800
814.104	15	1	15	320	4,800
814.106	10	1	10	120	1,200
814.108	12	1	12	80	960
814.110(a)	1	1	1	80	80
814.112(b)	1	1	1	8	8
814.116(b)	12	1	12	8	96
814.118(d)	1	1	1	8	8
814.120(b)	10	1	10	200	2,000
814.124(b)	2	1	2	2	4
814.126(b)(I)	2	1	2	120	240
Total					10,196

There are no capital costs or operating and maintenance costs associated with this collection.

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN

21 CFR Section	No. of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours per Recordkeeper	Total Hours
814.126(b)(ii)	12	1	12	2	24
Total					24

There are no capital costs or operating and maintenance costs associated with this collection.

Dated: October 24, 1996.

William B. Schultz,

Deputy Commissioner for Policy.

[FR Doc. 96–27737 Filed 10–24–96; 3:21 pm]

BILLING CODE 4160–01–F

[Docket No. 96N–0325]

Agency Information Collection Activities; Submission for OMB Review; Comment Request

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.

DATES: Submit written comments on the collection of information by November 29, 1996.

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, Attention: Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, rm. 16B–19, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: In compliance with section 3507 of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507), FDA has submitted the following proposed collection of information to OMB for review and clearance: Food Canning Establishment Registration, Process Filing and Recordkeeping for Acidified Foods and Thermally Processed Low-Acid Foods in Hermetically Sealed Containers—(21 CFR 108.25(c)(1) and (c)(2), (d), (e), (g); 108.35(c)(1), (c)(2), (d), (e), (f), (h); 113.60(c); 113.83; 113.87; 113.89; 113.100; 114.80(b); 114.89; 114.100(a) through (d)) (OMB Control Number 0910–0037—Reinstatement).

Under the Federal Food, Drug, and Cosmetic Act (the act), FDA is authorized to prevent the interstate distribution of food products that may be injurious to health or that are otherwise adulterated, as defined in