acquire voting shares of Farmers State Bankshares, Inc., Cheyenne, Wyoming, and thereby indirectly acquire Wyoming Bank & Trust, Cheyenne, Wyoming.

Board of Governors of the Federal Reserve System, November 20, 1997.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 97–30967 Filed 11–25–97; 8:45 am]

BILLING CODE 6210–01–F

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

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

TIME AND DATE: 11:00 a.m., Monday, December 1, 1997.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C 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: Joseph R. Coyne, 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.bog.frb.fed.us for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: November 21, 1997.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 97–31151 Filed 11–21–97; 4:09 pm]

BILLING CODE 6210–01–P

GENERAL SERVICES ADMINISTRATION

Performance Review Board; Membership; Senior Executive Service

AGENCY: General Services Administration.

ACTION: Notice.

SUMMARY: Notice is hereby given of the names of the members of the Performance Review Board.

FOR FURTHER INFORMATION CONTACT: Gail T. Lovelace, Director of Human Resources, General Services Administration, 1800 F Street, N.W., Washington, DC 20405, (202) 501–0398.

SUPPLEMENTARY INFORMATION: Section 4313(c) (1) through (5) of Title 5 U.S.C. requires each agency to establish in accordance with regulations prescribed by the Office of Personnel Management, one or more Performance Review Board(s). The Board(s) shall review the performance rating of each senior executive's performance by the supervisor, along with any recommendation to the appointing authority relative to the performance of the senior executive.

Members of the Review Board are:

1. Thurman M. Davis, Sr. (Chairperson), Deputy Administrator.
2. Martha N. Johnson, Chief of Staff.
3. Dennis J. Fisher, Commissioner, Federal Technology Service.
4. Robert A. Peck, Commissioner, Public Buildings Service.
5. Frank P. Pugliese, Commissioner, Federal Supply Service.
6. G. Martin Wagner, Associate Administrator for Governmentwide Policy.

Dated: November 20, 1997.

Gail T. Lovelace,

Director of Human Resources.

[FR Doc. 97–31038 Filed 11–25–97; 8:45 am]

BILLING CODE 6820–BR–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0472]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each reinstatement of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on requirements for filing a petition for administrative stay of action.

DATES: Submit written comments on the collection of information by January 26, 1998.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502 (3) and 5 CFR 1320.3 (c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506 (c) (2) (A) of the PRA (44 U.S.C. 3506 (c) (2) (A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Petition For Administrative Stay of Action—21 CFR Part 10.35 (OMB Control Number 0910—0194)—Reinstatement

Section 10.35 (21 CFR 10.35), issued under the authority of section 701(a) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 371(a)), sets forth the format and procedures by which an interested person may file a petition for an administrative stay of action.

Respondents to this information collection are interested persons who choose to file a petition for an administrative stay of action. Such a petition must: (1) Identify the decision involved; (2) state the action requested—including the length of time for which a stay is requested; and (3) include a statement of the factual and legal grounds on which the interested person relies in seeking the stay. The information provided in the petition is used by the agency to determine whether the requested stay should be granted.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
10.35	7	1	7	100	700

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

The burden estimate for this collection of information is based on FDA's experience with petitions for administrative stay of action over the past 3 years. Agency personnel responsible for processing the filing of petitions for administrative stays of action estimate that seven such petitions are received by the agency annually, with each requiring approximately 100 hours of preparation time.

Dated: November 19, 1997

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 97–30982 Filed 11–25–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 91N–0396]

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 (the PRA).

DATES: Submit written comments on the collection of information by December 26, 1997.

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, Attn: Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

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

Reports of Corrections and Removals for Manufacturers, Importers, and Distributors of Medical Devices (21 CFR 806.10 and 806.20).

In a final rule published in the **Federal Register** of May 19, 1997 (62 FR 27183), FDA issued regulations requiring that manufacturers, importers, and distributors of medical devices report promptly to FDA any corrections or removals of a device undertaken to reduce a risk to health posed by the device or to remedy a violation of the Federal Food, Drug, and Cosmetic Act (the act) that could present a risk to health. The collection of this information is required by section 519(f) of the act (21 U.S.C. 360i(f)). These regulations will help FDA to protect the public health by ensuring that the agency has current and complete information regarding those actions taken to reduce risks to health caused by devices. Reports of such actions will improve the agency's ability to evaluate device-related problems and to take prompt action against potentially dangerous devices.

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
806.10	880	1	880	10	8,800

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN[1]

21 CFR Section	No. of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours per Recordkeeper	Total Hours
806.20	440	1	440	10	4,400

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