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 September 16, 1985.

A. **Federal Reserve Bank of Atlanta** (Robert E. Heck, Vice President) 104 Marietta Street NW., Atlanta, Georgia 30303:

1. *Hub Bancshares, Inc.*, Lafayette, Louisiana; to engage *de novo* through its subsidiary, Hub Services, Inc., Lafayette, Louisiana, in real estate appraising as permitted by § 225.25(b)(13) of Regulation Y. This activity will be conducted in the south central Louisiana area.

2. *The First Bankers Corp. of Florida*, Pompano Beach, Florida; to engage *de novo* through its subsidiary, The First Bankers Management Group, Inc., Pompano Beach, Florida, in management and consulting advice to bank and nonbank depository institutions. These activities would be conducted in the central and southern Florida area.

Board of Governors of the Federal Reserve System, August 19, 1985.

James McAfee,

Associate Secretary of the Board.

[FR Doc. 85-20177 Filed 8-22-85; 8:45 am]

BILLING CODE 6210-01-M

Southwest First Community, Inc.; Application To Engage de Novo in Permissible Nonbanking Activities

The company listed in this notice has filed an application under § 225.23(a)(1) of the Board's Regulation Y (12 CFR 225.23(a)(1)) for the Board's approval under section 4(c)(8) of the Bank Holding Company Act (12 U.S.C. 1843(c)(8)) and §225.21(a) of Regulation Y (12 CFR 225.21(a)) to commence or to engage *de novo*, either directly or through a subsidiary, in a nonbanking activity that is listed in § 225.25 of Regulation Y as closely related to banking and permissible for bank holding companies. Unless otherwise noted, such activities will be conducted throughout the United States.

The application is available for immediate inspection at the Federal Reserve Bank 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 question 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." 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 application must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than September 13, 1985.

A. **Federal Reserve Bank of Dallas** (Anthony J. Montelaro, Vice President) 400 South Akard Street, Dallas, Texas 75222:

1. *Southwest First Community, Inc.*, Beeville, Texas; to engage *de novo* through its subsidiary, Southwest First Community Data, Inc., Beeville, Texas, in providing data processing and data transmission services; facilities and data bases (including data processing and data transmission hardware; software or operating personal) or access to such services, facilities or data bases.

Board of Governors of the Federal Reserve System, August 19, 1985.

James McAfee,

Associate Secretary of the Board.

[FR Doc. 85-20178 Filed 8-22-85; 8:45 am]

BILLING CODE 6210-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Commission on the Evaluation of Pain; Meeting

AGENCY: Department of Health and Human Services.

ACTION: Notice of Meeting.

SUMMARY: This notice announces the schedule and proposed agenda of the forthcoming meeting of the Commission on the Evaluation of Pain (the Commission). This notice also describes the purpose, structure, and termination date of the Commission.

DATES:

General session—September 5, 1985, 8:30 a.m. to 5:00 p.m.

General session—September 6, 1985, 8:30 a.m. to 5:00 p.m.

ADDRESS: National Academy of Sciences, 2101 Constitution Avenue NW., Washington, DC 20037.

FOR FURTHER INFORMATION CONTACT: Nancy Dapper, Executive Director, Commission on the Evaluation of Pain, Room 118, Altmeyer Building, 6401 Security Boulevard, Baltimore, Maryland 21235, (301) 597-1597.

SUPPLEMENTARY INFORMATION: The Commission is established and governed by the provisions of section 3(b) (1) through (6) of the Social Security Disability Benefits Reform Act of 1984 (Pub. L. 98-460). The purpose of the Commission is to conduct a study concerning the evaluation of pain in determining under titles II and XVI of the Social Security Act whether an individual is under a disability. The study is to be conducted in consultation with the National Academy of Sciences.

The study will consist of expert testimony and a review of research data regarding how pain should be considered in making disability determinations under titles II and XVI. The Commission may engage technical assistance in order to carry out its function.

The Commission is to submit a report, consisting of the findings of the study and any recommendations, to the Secretary of Health and Human Services (the Secretary) who in turn is to submit the report to the Committee on Ways and Means of the House of Representatives and to the Committee on Finance of the Senate.

The statute provides that the Commission terminate on December 31, 1985. This is also the deadline for the Secretary to submit the report.

The Secretary has appointed the members of the Commission in accordance with the provisions of the statute. This notice announces the third working meeting of the Commission. The Commission is chaired by Kathleen M. Foley, M.D.

This meeting is open to the public. Anyone wishing to submit his or her views for consideration by the Commission should send them to the Executive Director of the Commission at her address shown above.

A transcript of the Commission meeting will be made available to the public on an at-cost-of-duplication basis. The transcript can be ordered from the Executive Director of the Commission.

Agenda

September 5, 1985

8:30 a.m.—General session of expert testimony and research data presentations.

5:00 p.m.—Adjourn general session.

September 6, 1985

8:30 a.m.—General session of (1) continued expert testimony and research data presentations, (2) discussion on the agenda, priorities, and date of the next meeting, and (3) writing assignments.

5:00 p.m.—Adjourn the meeting.

Dated: August 19, 1985.

Nancy J. Dapper,

Executive Director, Commission on the Evaluation of Pain.

[FR Doc. 85-20227 Filed 8-22-85; 8:45 am]

BILLING CODE 4190-11-M

Food and Drug Administration

Ad Hoc Advisory Committee on Hypersensitivity to Food Constituents; Meetings

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: This notice announces forthcoming meetings of public advisory committees of the Food and Drug Administration (FDA). This notice also summarizes the procedures for the meetings and methods by which interested persons may participate in open public hearings before FDA's advisory committees.

Meetings: The following advisory committee meetings are announced:

Ad Hoc Advisory Committee on Hypersensitivity to Food Constituents

Date, time, and place. September 12 and 13, 9 a.m., Auditorium, 330 Independence Ave. SW., Washington, DC.

Type of meeting and contact person. Open committee discussion, September 12, 9 a.m. to 12 m.; open public hearing, 1:30 p.m. to 2:30 p.m.; open committee discussion, 2:30 to 4 p.m., September 13, 9 a.m. to 12 m.; Mary C. Custer, Center for Food Safety and Applied Nutrition (HFF-334), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-426-9463.

General function of the committee. The committee will review and evalaute available information relevant to adverse reactions in humans associated with the use of food constituents.

Agenda—Open public hearing. Interested persons requesting to present data, information, or views, orally or in writing, on issues pending before the committee should communicate with the committee contact person.

Open committee discussion. The committee will hear presentations on the terminology associated with adverse reactions to foods, a clinical approach to adverse reactions to foods, and patient strategy for coping with adverse reaction to foods.

FDA public advisory committee meetings may have as many as four separable portions: (1) An open public hearing, (2) an open committee discussion, (3) a closed presentation of data, and (4) a closed committee deliberation. Every advisory committee meeting shall have an open public hearing portion. Whether or not it also includes any of the other three portions will depend upon the specific meeting involved. There are no closed portions for the meetings announced in this notice. The dates and times reserved for the open portions of each committee meeting are listed above.

The open public hearing portion of each meeting shall be at least 1 hour long unless public participation does not last that long. It is emphasized, however, that the 1 hour time limit for an open public hearing represents a minimum rather than a maximum time for public participation, and an open public hearing may last for whatever longer period the committee chairman determines will facilitate the committee's work.

Public hearings are subject to FDA's guideline (Subpart C of 21 CFR Part 10) concerning the policy and procedures to expedite electronic media coverage of FDA's public administrative proceedings, including hearings before public advisory committees under 21 CFR Part 14. Under 21 CFR 10.205, representatives of the electronic media may be permitted, subject to certain limitations, to videotape, film, or otherwise record FDA's public administrative proceedings, including presentations by participants.

Meetings of advisory committees shall be conducted, insofar as is practical, in accordance with the agenda published in this **Federal Register** notice. Changes in the agenda will be announced at the beginning of the open portion of a meeting.

Any interested person who wishes to be assured of the right to make an oral presentation at the open public hearing portion of a meeting shall inform the contact person listed above, either orally or in writing, prior to the meeting. Any person attending the hearing who does not in advance of the meeting request an opportunity to speak will be allowed to make an oral presentation at the hearing's conclusion, if time permits, at the chairman's discretion.

Persons interested in specific agenda items to be discussed in open session may ascertain from the contact person the approximate time of discussion.

A list of committee members and summary minutes of meetings may be requested from the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857, between the hours of 9 a.m. and 4 p.m., Monday through Friday.

This notice is issued under section 10(a) (1) and (2) of the Federal Advisory Committee Act (Pub. L. 92-463, 86 Stat. 770-776 (5 U.S.C. App. I)), and FDA's regulations (21 CFR Part 14) on advisory committees.

Dated: August 19, 1985.

Mervin H. Shumate,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 85-20166 Filed 8-20-85; 10:07 am]

BILLING CODE 4160-01-M

[Docket No. 85F-0346]

Foodways National, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Foodways National, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of aspartame in frozen cheesecake, fruit, and fruit toppings.

FOR FURTHER INFORMATION CONTACT: Anthony P. Brunetti, Center for Food Safety and Applied Nutrition (HFF-334), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-426-5487.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 5A3874) has been filed by Foodways National, Inc., P.O. Box 44, Boise, ID 83707, proposing that § 172.804 *Aspartame* (21 CFR 172.804) be amended to provide for the safe use of aspartame to sweeten frozen cheesecake, fruit, and fruit toppings.

The potential environmental impact of this action is being reviewed. If the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's

finding of no significant impact and the evidence supporting that finding will be published with the regulation in the Federal Register in accordance with 21 CFR 25.40(c), as published in the Federal Register of April 26, 1985 (50 FR 16636).

Dated: August 14, 1985.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 85-20161 Filed 8-22-85; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 85F-0345]

Pfizer, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Pfizer, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of aspartame in frozen desserts.

FOR FURTHER INFORMATION CONTACT: Anthony P. Brunetti, Center for Food Safety and Applied Nutrition (HFF-334), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-426-5487.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 5A3861) has been filed by Pfizer Central Research, Pfizer, Inc., 235 East 42nd Street, New York, NY 10017, proposing that § 172.804(c) (21 CFR 172.804(c)) be amended to provide for the safe use of aspartame to sweeten frozen desserts, where standards of identity do not preclude this use.

The potential environmental impact of this action is being reviewed. If the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the Federal Register in accordance with 21 CFR 25.40(c), as published in the Federal Register of April 26, 1985 (50 FR 16636).

Dated: August 15, 1985.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 85-20160 Filed 8-22-85; 8:45 am]

BILLING CODE 4160-01-M

Health Resources and Services Administration

Application Announcement for Grants for Predoctoral Training in Family Medicine

The Bureau of of Health Professions, Health Resources and Services Administration, announces that applications for Fiscal Year 1986 Grants for Predoctoral Training in Family Medicine are being accepted under the authority of section 786(a) of the Public Health Service Act, as amended.

Applicants should be advised that this application announcement is a contingency action being taken to ensure that should funds become available for this purpose, they can be awarded in a timely fashion consistent with the needs of the programs as well as to provide for even distribution of funds throughout the fiscal year. The Administration's budget request for Fiscal Year 1986 does not include funding for this program. This notice regarding applications does not reflect any change in this policy.

In addition, programmatic changes may result from currently ending legislative action. Should such changes be necessary, all applicants will be notified at a later date.

Section 786(a) of the Public Health Service Act authorizes the award of grant to assist in meeting the cost of planning, developing and operating or participating in approved predoctoral training programs in the field of family medicine. Grants may include support for the program only or support for both the program and the trainees.

To receive support, programs must meet the requirements of regulations as set forth in 42 CFR Part 57, Subpart Q. Eligible applicants are accredited public or nonprofit private schools of medicine or osteopathic medicine. In accordance with § 57.1605(b)(2), a funding preference will be accorded approved applications for projects in which:

1. Substantial training experience is in settings which exemplify interdependent utilization of physicians and physician assistants or nurse practitioners; and/or

2. Substantial portions of the training program are conducted in a primary medical care manpower shortage area which is a part of a health manpower shortage area(s) designated under Section 332 of the PHS Act, or, in an Area Health Education Center funded, at least in part, under Section 781 of the Act.

Requests for application materials and questions regarding grants policy should be directed to: Grants Management Officer (D-15), Bureau of Health Professions, Health Resources and Services Administration, Parklawn Building, Room 4C-16, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone: (301) 443-6960.

Should additional programmatic information be required, please contact: Multidisciplinary Resources Development Branch, Division of Medicine, Bureau of Health Professions, Health Resources and Services Administration, Parklawn Building, Room 8C-22, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone: (301) 443-3614.

The deadline date for receipt of applications is September 30, 1985. Applications shall be considered as meeting the deadline date if they are either:

(1) *Received* on or before the deadline date, or

(2) *Postmarked* on or before the deadline and received in time for submission to the independent review group. A legibly dated receipt from a commercial carrier or U.S. Postal Service will be accepted in lieu of a postmark. Private metered postmarks shall not be acceptable as proof of timely mailing.

This program is listed at 13.986 in the *Catalog of Federal Domestic Assistance.* It is not subject to the provisions of Executive Order 12372, Intergovernmental Review of Federal Programs or 45 CFR Part 100.

Dated: August 19, 1985.

John H. Kelso,

Acting Administrator.

[FR Doc. 85-20168 Filed 8-22-85; 8:45 am]

BILLING CODE 4160-16-M

Public Health Service

Recovery of Funds in Transactions Affecting Ownership of Federally Assisted Health Professions and Nurse Teaching Facilities

AGENCY: Health Resources and Services Administration (HRSA); Public Health Service, HHS.

ACTION: Statement of General Policy.

SUMMARY: HRSA announces its recovery policy regarding purported leases and sales of (1) health professions teaching facilities or affiliated hospitals constructed with grant funds under Part B of Title VII of the Public Health Service Act, and (2) nurse teaching facilities constructed with grant funds under Title VIII of the Act.