indicated or the offices of the Board of Governors not later than June 16, 2008.

A. Federal Reserve Bank of Richmond (A. Linwood Gill, III, Vice President) 701 East Byrd Street, Richmond, Virginia 23261–4528:

1. Eagle Bancorp, Inc.; to acquire 100 percent of the voting shares of Fidelity & Trust Financial Corporation, and thereby indirectly acquire Fidelity & Trust Bank, all of Bethesda, Maryland.

Board of Governors of the Federal Reserve System, May 19, 2008.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E8–11463 Filed 5–21–08; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

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, including the companies listed below, that engages either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.28 of Regulation Y (12 CFR 225.28) 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. The notice also will 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. 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 the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than June 6, 2008.

A. Federal Reserve Bank of Dallas (W. Arthur Tribble, Vice President) 2200 North Pearl Street, Dallas, Texas 75201–2272:

1. McCamey Financial Corporation, Odessa, Texas, and McCamey Financial Delaware Corporation, Dover, Delaware, through its subsidiary, Security State Bank, Odessa, Texas, to acquire 70 percent of the voting shares of Venture Finance LLC, Midland, Texas, and thereby engage in lending activities pursuant to section 225.28(b)(1) of Regulation Y.

Board of Governors of the Federal Reserve System, May 19, 2008.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E8–11462 Filed 5–21–08; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Health Marketing

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meetings:

Name: Board of Scientific Counselors, National Center for Health Marketing (BSC, NCHM).

Times and Dates: 10 a.m.–5 p.m., June 5, 2008. 8:30 a.m.–12 p.m., June 6, 2008.

Place: Auditorium A, Global Communications Center, Building 19, 1600 Clifton Road, N.E., Atlanta, Georgia, 30333.

Status: Open to the public, limited only by the space available.

Please Note: Due to current security measures, a valid government issued identification card with photo is required for admittance into the Roybal facility. Non-U.S. citizens wishing to attend should contact Dionne Mason; Telephone, (404) 498–2314. The deadline for notification of attendance is May 22, 2008.

Purpose: The board provides advice to the Secretary, Department of Health and Human Services; and the Director, Centers for Disease Control and Prevention, on strategies and goals for the programs and research within the national center; conducts peer review of scientific programs; and monitors the overall strategic direction and focus of the national center. The board also performs second-level peer review of applications for grants-in-aid for research and research training activities, cooperative agreements, and research contract proposals relating to the broad areas within the national center.

Matters to be Discussed: The agenda will include a general overview of the NCHM and discussions related to the Center's role in preparedness, response and recovery with regards to an outbreak of pandemic influenza.

Agenda items are subject to change as priorities dictate.

Contact for More Information: Dionne R. Mason, Committee Management Specialist, NCHM, CDC, 1600 Clifton Road, NE., Mail Stop E–21, Atlanta, Georgia 30333; Telephone, (404) 498–2314; Fax, (404) 498–2221.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: May 9, 2008.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E8–11448 Filed 5–21–08; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0286]

Agency Information Collection Activities; Proposed Collection; Comment Request; Survey to Evaluate FDA's Food Defense Awareness Initiative ALERT

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on a survey of food industry supervisory employees about their awareness and perceptions of FDA's Food Defense Awareness Initiative ALERT.

DATES: Submit written or electronic comments on the collection of information by July 21, 2008.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food

and Drug Administration, 5600 Fishers Lane, 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 before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (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.

Survey to Evaluate FDA's Food Defense Awareness Initiative ALERT

In July 2006, FDA announced its Food Defense Awareness Initiative, called ALERT (the letters stand for the five key components of the initiative: (assure, look, employees, report, and threat). The ALERT initiative is intended to raise the awareness of State and local government agencies and the food industry regarding food defense issues. ALERT identifies five key points that industry and businesses can use to decrease the risk of intentional food contamination at their facility. The ALERT Web-based training module and more information on ALERT are available at *www.cfsan.fda.gov/~dms/defterr.html*.

Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393 (b)(2)), FDA is authorized to conduct research relating to foods and to conduct educational and public information programs relating to the safety of the nation's food supply. Under this authority, FDA is planning to conduct a survey of first line supervisors working in a range of capacities in the food industry about their awareness and perceptions of the agency's ALERT initiative and the ALERT initiative informational materials. The purpose of the survey is to help FDA evaluate ALERT informational materials and to gauge whether the materials succeed in informing food industry supervisory employees about the risk of intentional food contamination and in motivating them to engage in protective behaviors. The survey results will be used to assess how knowledge and awareness, threat perceptions, attitudes, norms, benefits and barriers affect the implementation of the ALERT initiative.

The data will be collected using a Web-based questionnaire. The survey will employ a stratified, cluster sampling design. Using industry networks and listings, we will randomly sample from databases of eight industry groups (regulators, growers, packers, processors, warehousers, transporters, retailers, and food service operators). We will stratify within groups by organization size (small, medium, and large) based on number of employees on the payroll, for a total random sample of 200 organizations. Participation in the survey is voluntary. Cognitive interviews and a pre-test will be conducted prior to fielding the survey.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Questionnaire	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours Per Response	Total Hours
Cognitive Interviews	10	10	10	1	10
Pre-tests	10	1	10	.4	4
Survey	200	1	200	.4	80
Total					94

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

FDA's burden estimate is based on prior experience with consumer surveys similar to this proposed survey.

Please note that on January 15, 2008, the FDA Division of Dockets Management Web site transitioned to the Federal Dockets Management System (FDMS). FDMS is a Government-wide, electronic docket management system. Electronic comments or submissions will be accepted by FDA only through FDMS at *http://www.regulations.gov*.

Dated: May 15, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8–11514 Filed 5–21–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–D–0293]

Draft Guidance for Industry: Considerations for Allogeneic Pancreatic Islet Cell Products; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the