Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the office of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than August 13, 2008.

- A. Federal Reserve Bank of Minneapolis (Jacqueline G. King, Community Affairs Officer) 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291:
- 1. David J. Schornack, and Denise N. Schornack, both of Perham, Minnesota; Daniel R. Welter, and Bonnie S. Welter, both of New York Mills, Minnesota, to acquire voting shares of Quality Bankshares, Inc., and thereby indirectly acquire voting shares of Quality Bank, both of Fingal, North Dakota.
- **B. Federal Reserve Bank of Kansas City** (Todd Offenbacker, Assistant Vice
  President) 1 Memorial Drive, Kansas
  City, Missouri 64198–0001:
- 1. Robyn E. Batson, Broken Bow, Oklahoma, as trustee of the Linda Lake Young Irrevocable Trust, the Lori Lee Young Irrevocable Trust, and the Robyn Elizabeth Batson Irrevocable Trust, to acquire voting shares of Southeastern Bancshares, Inc., and thereby indirectly acquire voting shares of 1st Bank & Trust, both in Broken Bow, Oklahoma.
- C. Federal Reserve Bank of Dallas (W. Arthur Tribble, Vice President) 2200 North Pearl Street, Dallas, Texas 75201– 2272.
- 1. John Mangan, San Antonio, Texas, as independent trustee of the LCK 1993 Trust, LCK Dynasty Trust, LCK Legacy Trust, the Katz Millennium Trust, and LCK Trust Number 2, to acquire voting shares of First Community Bancshares, Inc., Houston, Texas, and thereby indirectly acquire voting shares of First National Bank Texas, Killeen, Texas, and Fort Hood National Bank, Fort Hood, Texas.

Board of Governors of the Federal Reserve System, July 24, 2008.

#### Robert deV. Frierson,

Deputy Secretary of the Board. [FR Doc. E8–17355 Filed 7–28–08; 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: 12:00 p.m., Monday, August 4, 2008.

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.

# **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 <a href="http://www.federalreserve.gov">http://www.federalreserve.gov</a> 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, July 25, 2008.

#### Robert deV. Frierson,

Deputy Secretary of the Board. [FR Doc. 08–1476 Filed 7–25–08; 2:14 pm] BILLING CODE 6210–01–S

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; American Health Information Community Consumer Empowerment Workgroup Meeting

**ACTION:** Announcement of meeting.

**SUMMARY:** This notice announces the 28th meeting of the American Health Information Community Consumer Empowerment Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. No. 92–463, 5 U.S.C., App.).

**DATES:** August 19, 2008, from 1 p.m. to 4 p.m. [Eastern]

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC

20201), Conference Room 1114. Please bring photo ID for entry to a Federal building.

## FOR FURTHER INFORMATION CONTACT:

http://www.hhs.gov/healthit/ahic/consumer/.

**SUPPLEMENTARY INFORMATION:** The Workgroup will continue its discussion on how to encourage the widespread adoption of a personal health record that is easy-to-use, portable, longitudinal, affordable, and consumercentered.

The meeting will be available via Web cast. For additional information, go to: http://www.hhs.gov/healthit/ahic/consumer/ce\_instruct.html.

### Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. E8–17297 Filed 7–28–08; 8:45 am] **BILLING CODE 4150–45–P** 

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; American Health Information Community Quality Workgroup Meeting

**ACTION:** Announcement of meeting.

**SUMMARY:** This notice announces the 19th meeting of the American Health Information Community Quality Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. No. 92–463, 5 U.S.C., App.).

**DATES:** August 5, 2008, from 1 p.m. to 4 p.m. [Eastern].

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 1114. Please use the C Street entrance closest to 3rd Street and bring photo ID for entry to a Federal building.

FOR FURTHER INFORMATION CONTACT: http://www.hhs.gov/healthit/ahic/qualitv/.

SUPPLEMENTARY INFORMATION: The Workgroup will continue its discussion on how health information technology can provide the data needed for the development of quality measures that are useful to patients and others in the health care industry, automate the measurement and reporting of a comprehensive current and future set of quality measures, and accelerate the use of clinical decision support that can improve performance on those quality measures.

The meeting will be available via Web cast. For additional information, go to: http://www.hhs.gov/healthit/ahic/ quality/quality\_instruct.html.

#### Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. E8-17298 Filed 7-28-08; 8:45 am] BILLING CODE 4150-45-P

#### DEPARTMENT OF HEALTH AND **HUMAN SERVICES**

Office of the National Coordinator for Health Information Technology; **American Health Information** Community Confidentiality, Privacy, & **Security Workgroup Meeting** 

**ACTION:** Announcement of meeting.

**SUMMARY:** This notice announces the 22nd meeting of the American Health Information Community Confidentiality, Privacy, & Security Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. No. 92-463, 5 U.S.C., App.).

**DATES:** August 21, 2008, from 1 p.m. to 5 p.m. [Eastern Time].

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 1114. Please use the C Street entrance closest to 3rd Street and bring photo ID for entry to a Federal building.

## FOR FURTHER INFORMATION CONACT:

http://www.hhs.gov/healthit/ahic/ confidentiality/.

SUPPLEMENTARY INFORMATION: The Workgroup Members will continue discussing and evaluating the confidentiality, privacy, and security protections and requirements for participants in electronic health information exchange environments.

The meeting will be available via Web cast. For additional information, go to: http://www.hhs.gov/healthit/ahic/ cps\_instruct.html.

# Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. E8-17313 Filed 7-28-08; 8:45 am]

BILLING CODE 4150-45-P

### **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Office of the National Coordinator for Health Information Technology; **American Health Information Community Population Health and Clinical Care Connections Workgroup** Meeting

**ACTION:** Announcement of meeting.

**SUMMARY:** This notice announces the 29th meeting of the American Health Information Community Population Health and Clinical Care Connections Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. 92-463, 5 U.S.C., App.).

**DATES:** August 20, 2008, from 2 p.m. to 5 p.m. [Eastern Time].

ADDRESSES: Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 1114. Please use the C Street entrance closest to 3rd Street and bring photo ID for entry to a Federal building.

# FOR FURTHER INFORMATION CONTACT:

http://www.hhs.gov/healthit/ahic/ population/.

SUPPLEMENTARY INFORMATION: The Workgroup will continue its discussion on how to facilitate the flow of reliable health information among population health and clinical care systems necessary to protect and improve the public's health. The meeting will be available via Web cast. For additional information, go to: http://www.hhs.gov/ healthit/ahic/population/ pop\_instruct.html.

## **Judith Sparrow**,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. E8-17314 Filed 7-28-08; 8:45 am] BILLING CODE 4150-45-P

# **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Food and Drug Administration

[Docket No. FDA-2008-N-0389]

**Food and Drug Administration** Amendments Act of 2007; Prohibition Against Food to Which Drugs or **Biological Products Have Been Added; Request for Comments** 

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA) is requesting

comments relevant to the implementation of section 912 of the Food and Drug Administration Amendments Act of 2007 (FDAAA). Section 912 of FDAAA establishes section 301(ll) in the Federal Food. Drug, and Cosmetic Act (the act), which prohibits the interstate shipment of certain foods to which an approved drug or a licensed biological product has been added. Section 301(ll) also prohibits the interstate shipment of foods containing an added drug or a biological product that has been the subject of substantial clinical investigations, the existence of which has been made public. FDA requests that interested persons submit data, information, and comments that will help provide a context for the agency's decisions on implementation of this provision. To encourage responsive comments, FDA is including a series of questions for interested persons to consider in preparing comments.

**DATES:** Submit written or electronic comments by October 27, 2008.

**ADDRESSES:** Submit written comments, data, and other information to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to www.regulations.gov.

#### FOR FURTHER INFORMATION CONTACT:

Catherine L. Copp, Center for Food Safety and Applied Nutrition (HFS-4), Food and Drug Administration, 301-436-1589, e-mail: catherine.copp@ fda.hhs.gov.

# SUPPLEMENTARY INFORMATION:

# I. Background

On September 27, 2007, the Food and Drug Administration Amendments Act of 2007 (Public Law 110-85) (FDAAA) was enacted. Section 912 of FDAAA establishes section 301(ll) in the Federal Food, Drug, and Cosmetic Act (the act), 21 U.S.C. 331(ll), which adds the following prohibited act to section 301.21 U.S.C. 331:

The introduction or delivery for introduction into interstate commerce of any food to which has been added a drug approved under section 505, a biological product licensed under section 351 of the Public Health Service Act, or a drug or a biological product for which substantial clinical investigations have been instituted and for which the existence of such investigations has been made public, unless--

(1) such drug or such biological product was marketed infood before any approval of the drug under section 505, beforelicensure of the biological product under such section 351, andbefore any substantial clinical