

Board of Governors of the Federal Reserve System, November 10, 2004.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 04-25454 Filed 11-16-04; 8:45 am]

BILLING CODE 6210-01-S

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisition of Shares of Bank or Bank Holding Companies; Correction

This notice corrects a notice (FR Doc. 04-24999) published on pages 65195 and 65196 of the issue for Wednesday, November 10, 2004.

Under the Federal Reserve Bank of St. Louis heading, the entry for Charles Keith Akin, is revised to read as follows:

A. Federal Reserve Bank of St. Louis (Randall C. Sumner, Vice President) 411 Locust Street, St. Louis, Missouri 63166-2034:

1. *Charles Keith Akin*, Clinton, Kentucky; and the Akin Control Group, which consists of Charles Keith Akin; Anita Akin; Burkley Investments, Inc.; Parkway Manor – KY; and Parkway Manor – TN, all of Clinton, Kentucky; and Bruce Akin, Paducah, Kentucky; to acquire additional voting shares of Purchase Area Bancorp, Bardwell, Kentucky, and thereby indirectly acquire voting shares of Bardwell Deposit Bank, Bardwell, Kentucky.

Comments on this application must be received by November 24, 2004.

Board of Governors of the Federal Reserve System, November 10, 2004.

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Deputy Secretary of the Board.

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FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR Part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The applications listed below, as well as other related filings required by the Board, are available for immediate

inspection at the Federal Reserve Bank indicated. The application also will 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 (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States. 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 each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than December 10, 2004.

A. Federal Reserve Bank of Chicago (Patrick Wilder, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. *QCR Holdings, Inc.*, Moline, Illinois; to acquire 100 percent of the voting shares of Rockford Bank and Trust Company, Rockford, Illinois (in organization).

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Deputy Secretary of the Board.

[FR Doc. 04-25453 Filed 11-16-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research And Quality

Notice of Meeting

In accordance with section 10(d) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2), announcement is made of a Health Care Policy and Research Special Emphasis Panel (SEP) meeting.

A Special Emphasis panel is a group of experts in fields related to health care research who are invited by the Agency for Healthcare Research and Quality (AHRQ), and agree to be available, to conduct on an as needed basis, scientific reviews of applications of AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in

particular review meetings which require their type of expertise.

Substantial segments of the upcoming SEP meeting listed below will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6). Peer Review of a Research Grant application (R03) will be discussed at this meeting. These discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure under the above-cited statutes.

SEP Meeting on: AHRQ Research Grant Application (R03).

Date: November 17, 2004 (Open November 17 from 1:30 p.m. to 1:45 p.m. and closed for the remainder of the teleconference meeting).

Place: John M. Eisenberg Building, AHRQ Conference Center, 540 Gaither Road, Rockville, Maryland 20850.

Contact Person: Anyone wishing to obtain a roster of members, agenda or minutes of the nonconfidential portions of this meeting should contact Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, AHRQ, 540 Gaither Road, Room 2038, Rockville, Maryland 20850, telephone (301) 427-1554.

Agenda items for this meeting are subject to change as priorities dictate.

This notice is being published less than 15 days prior to the November 17 meeting, due to the time constraints of reviews and funding cycles.

Dated: November 8, 2004.

Carolyn M. Clancy, M.D.,
Director.

[FR Doc. 04-25474 Filed 11-16-04; 8:45 am]

BILLING CODE 4160-90-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004D-0499]

Compliance Policy Guide; Radiofrequency Identification Feasibility Studies and Pilot Programs for Drugs; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a new compliance policy guide (CPG) Sec. 400.210 entitled

“Radiofrequency Identification Feasibility Studies and Pilot Programs for Drugs.” The CPG describes the agency’s intent to exercise enforcement discretion, until December 31, 2007, concerning certain regulatory requirements to facilitate the performance of feasibility studies and pilot programs involving Radiofrequency Identification (RFID) tags for drugs. The goal of the CPG is to allow industry to gain experience with the use of RFID technology to ensure the long-term safety and integrity of the U.S. drug supply.

DATES: You may submit written or electronic comments at any time.

ADDRESSES: Submit written requests for single copies of the guidance to the Division of Compliance Policy (HFC-230), Office of Enforcement, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857. Send one self-addressed adhesive label to assist that office in processing your request or include a fax number to which the guidance may be sent.

Submit written comments on the guidance to the Division of Dockets Management, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.fda.gov/dockets/ecomments>. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

FOR FURTHER INFORMATION CONTACT: Paul Rudolf, Office of Policy (HF-11), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-3360.

SUPPLEMENTARY INFORMATION:

I. Background

On February 18, 2004, FDA published a report entitled “Combating Counterfeit Drugs” which is available on the FDA Web site at <http://www.fda.gov/oc/initiatives/counterfeit>. In that report the agency identified RFID technology as the cornerstone in the fight against counterfeit drugs and announced our intention to facilitate the adoption of RFID technology by participants in the pharmaceutical supply chain. We also stated that widespread adoption of RFID technology was feasible by 2007.

Recently, FDA has received inquiries focusing on whether certain regulatory requirements, including those related to labeling, electronic records, and product quality, apply to pharmaceutical manufacturers, repackagers, relabelers, distributors, retailers, or others who participate in feasibility studies and pilot programs (collectively “a study” or “studies”) using RFID tags for drugs. This CPG describes how we intend to

exercise our enforcement discretion regarding such studies. The exercise of such enforcement discretion expires on December 31, 2007. The goal of this CPG is to facilitate the performance of RFID studies and allow industry to gain experience with the use of RFID.

FDA is issuing this document as a level 1 guidance consistent with FDA’s good guidance practices regulation (§ 10.115 (21 CFR 10.115)). The new CPG Sec. 400.210 is being implemented immediately without prior public comment under § 10.115(g)(2), because the agency has determined that prior public participation is not feasible or appropriate, but comments are welcome at any time. The agency also thinks that use of RFID technology is critical to ensuring the long-term safety and integrity of the U.S. drug supply and immediate guidance is needed to facilitate studies of RFID.

II. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments on the guidance document. Submit two copies of written comments, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. The guidance and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

III. Electronic Access

An electronic version of this guidance is available on the Internet at <http://www.fda.gov/ora> under “Compliance Reference.”

Dated: November 10, 2004.

John Taylor,

Associate Commissioner for Regulatory Affairs.

[FR Doc. 04-25527 Filed 11-15-04; 9:19 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HOMELAND SECURITY

Federal Law Enforcement Training Center

Notice of meeting

AGENCY: Federal Law Enforcement Training Center (FLETC), Department of Homeland Security.

ACTION: Notice of Meeting.

SUMMARY: The Advisory Committee to the National Center for State and Local Law Enforcement Training (National Center) at the Federal Law Enforcement

Training Center will meet on December 1, 2004, beginning at 8 a.m.

ADDRESSES: Federal Law Enforcement Training Center, 1131 Chapel Crossing Road, Glynco, GA 31524.

FOR FURTHER INFORMATION CONTACT:

Reba Fischer, Designated Federal Officer, National Center for State and Local Law Enforcement Training, Federal Law Enforcement Training Center, Glynco, GA 31524, 912-267-2343, reba.fischer@dhs.gov.

SUPPLEMENTARY INFORMATION: The agenda for this meeting includes remarks by the Committee Co-Chairs, Randy Beardsworth, Director of Operations, Border and Transportation Security, Department of Homeland Security, and Deborah Daniels, Assistant Attorney General, Office of Justice Programs, Department of Justice; an update on current training initiatives of the National Center; and planning of strategic goals. This meeting is open to the public. Anyone desiring to attend must contact Reba Fischer, the Designated Federal Officer, no later than November 20, 2004, at (912) 267-2343, to arrange clearance. This meeting was originally scheduled for September 14, 2004, but was cancelled due to Hurricane Ivan.

Dated: November 5, 2004.

Stanley Moran,

Director, National Center for State and Local Law Enforcement Training.

[FR Doc. 04-25545 Filed 11-12-04; 4:32 pm]

BILLING CODE 4810-32-U

DEPARTMENT OF HOMELAND SECURITY

Citizenship and Immigration Services Bureau

[CIS No. 2331-04]

RIN 1615-ZA68

Extension of Honduras for Temporary Protected Status; Correction

AGENCY: Citizenship and Immigration Services, Department of Homeland Security.

ACTION: Notice of correction.

SUMMARY: U.S. Citizenship and Immigration Services (USCIS) is correcting a notice that was published in the **Federal Register** on November 3, 2004 at 69 FR 64084 which announced the extension of the designation of Honduras for Temporary Protected Status (TPS). In the supplemental information to the notice, USCIS inadvertently misstated that only Form I-821 with Revision Date 7/30/04 will