EXPORT-IMPORT BANK OF THE UNITED STATES

Sunshine Act Meeting

AGENCY: Export-Import Bank of the United States.

ACTION: Cancellation of a Government in the Sunshine Meeting.

ORIGINAL TIME AND PLACE: Thursday, April 27, 2006 at 9:30 a.m.

PLACE: Room 1143, 811 Vermont Avenue, NW., Washington, DC 20571.

The Export-Import Bank of the United States has cancelled the Government in the Sunshine meeting which was scheduled for April 27, 2006. The Bank will reschedule this meeting at a future date. Earlier announcement of this cancellation was not possible.

FOR FURTHER INFORMATION CONTACT: For further information, contact: Office of the Secretary, 811 Vermont Avenue, NW., Washington, DC 20571 (Tele. No. 202–565–3957).

Howard A. Schweitzer,

General Counsel (Acting).

[FR Doc. 06–4101 Filed 4–26–06; 4:08 am]

BILLING CODE 6690–01–M

FEDERAL RESERVE SYSTEM

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

The notificants listed below have applied under the Change in Bank 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 May 16, 2006.

A. Federal Reserve Bank of Kansas City (Donna J. Ward, Assistant Vice President) 925 Grand Avenue, Kansas City, Missouri 64198-0001:

1. Biegert Family Trust, Laramie, Wyoming, its trustees, Larry R. Cox; Henderson, Nebraska, Judith Ackland, Geneva, Nebraska, and Larry R. Cox, individually; Charles Flaming, individually, and as owner of Sadle Cattle Company, Inc., both of Paxton, Nebraska; Alan Janzen, Christopher Vanderneck, Matthew D. Siebert, Fredrick Regier, Arvid Janzen, and Brian Janzen, all of Henderson, Nebraska; Ronald Preheim, Aurora, Nebraska; Jeff Pribbeno, Imperial, Nebraska; and Wesley Kroeker, Enid, Oklahoma; and thereby indirectly acquire shares of Henderson State Company, Henderson, Nebraska, of Henderson State Bank, Henderson, Nebraska.

Board of Governors of the Federal Reserve System, April 26, 2006.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E6–6530 Filed 4–28–06; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Federal Open Market Committee; Domestic Policy Directive of March 27 and 28, 2006

In accordance with § 271.25 of its rules regarding availability of information (12 CFR part 271), there is set forth below the domestic policy directive issued by the Federal Open Market Committee at its meeting held on March 27 and 28, 2006.[1]

The Federal Open Market Committee seeks monetary and financial conditions that will foster price stability and promote sustainable growth in output. To further its long-run objectives, the Committee in the immediate future seeks conditions in reserve markets consistent with increasing the federal funds rate to an average of around 4¾ percent.

The vote encompassed approval of the paragraph below for inclusion in the statement to be released shortly after the meeting:

''The Committee judges that some further policy firming may be needed to keep the risks to the attainment of both sustainable economic growth and price stability roughly in balance. In any event, the Committee will respond to changes in economic prospects as needed to foster these objectives.''

[1] Copies of the Minutes of the Federal Open Market Committee Meeting on March 27 and 28, 2006, which includes the domestic policy directive issued at the meeting, are available upon request to the Board of Governors of the Federal Reserve System, Washington, DC 20551. The minutes are published in the Federal Reserve Bulletin and in the Board's annual report.

By order of the Federal Open Market Committee, April 20, 2006.

Vincent R. Reinhart,

Secretary, Federal Open Market Committee.

[FR Doc. E6–6492 Filed 4–28–06; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–06–0222]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an e-mail to *omb@cdc.gov*.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

Questionnaire Design Research Laboratory (QDRL) 2007–2009, (OMB No. 0920–0222)—Extension—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Questionnaire Design Research Laboratory (QDRL) conducts questionnaire pre-testing and evaluation activities for CDC surveys (such as the NCHS National Health Interview

Survey, OMB No. 0920–0214) and other federally sponsored surveys. The QDRL conducts cognitive interviews, focus groups, mini field-pretests, and experimental research in laboratory and field settings, both for applied questionnaire evaluation and more basic research on response errors in surveys. The most common questionnaire evaluation method is the cognitive interview. In a cognitive interview, a questionnaire design specialist interviews a volunteer participant. The interviewer administers the draft survey questions as written, but also probes the participant in depth about interpretations of questions, recall processes used to answer them, and adequacy of response categories to express answers, while noting points of confusion and errors in responding. Interviews are generally conducted in small rounds of 10–15 interviews; ideally, the questionnaire is re-worked between rounds and revisions are tested iteratively until interviews yield relatively few new insights. When possible, cognitive interviews are conducted in the survey's intended mode of administration. For example, when testing telephone survey questionnaires, participants often respond to the questions via a telephone in a laboratory room. Under this condition, the participant answers without face-to-face interaction. QDRL staff watch for response difficulties from an observation room, and then conduct a face-to-face debriefing with in-depth probes. Cognitive interviewing provides useful data on questionnaire performance at minimal cost and respondent burden. Similar methodology has been adopted by other federal agencies, as well as by academic and commercial survey organizations. NCHS is requesting 3 years of OMB Clearance for the project. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN

Respondents	Number of respondents per year	Number of responses/ respondent	Avg. burden response (in hours)	Total burden hours
2007 test volunteers	500	1	1.2	600

Dated: April 25, 2006.

Joan F. Karr,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E6–6501 Filed 4–28–06; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0273] (formerly 03N–0273)

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Research Study Complaint Form

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Research Study Complaint Form'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 16, 2005 (70 FR 74817), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0579. The approval expires on March 31, 2009. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: April 24, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–6457 Filed 4–28–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0166]

Agency Emergency Processing Under the Office of Management and Budget Review; MedWatch—The Food and Drug Administration Safety Information and Adverse Event Reporting Program; Proposal to Survey MedWatch Partners Organizations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for emergency processing under the Paperwork Reduction Act of 1995 (the PRA). This notice solicits comments on a proposal for the MedWatch program to deploy and conduct a web-based customer satisfaction survey of certain health care professional trade and specialty organizations that voluntarily have chosen to participate in the FDA MedWatch's Partners program. The survey will solicit information about the utility of the FDA MedWatch safety alerts and monthly safety labeling changes that are posted on the MedWatch Web site and disseminated to partner organizations for sharing with members of the organizations.

DATES: Fax written comments on the collection of information by May 31, 2006. FDA is requesting approval of this emergency processing by May 31, 2006.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Fumie Yokota, Desk Officer for FDA, Fax: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Karen Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: FDA has requested emergency processing of this