1. Memorandum item 1, collects information on the number and amount of deposit accounts of (a) $100,000 or less and (b) more than $100,000. This information provides the basis for calculating ''simple estimates'' of the amount of insured and uninsured deposits. The captions for these memorandum items explicitly refer to $100,000, which is the current deposit insurance limit. Given the purpose of these memorandum items, the dollar amount cited in the caption will be changed if the deposit insurance limit were to change. The revision will ensure that such a change occurs automatically as a function of the deposit insurance limit in effect on the report date.

Schedule S—Securitization and Asset Sale Activities

Splitting item 2.b., ''Standby letters of credit, subordinated securities, and other enhancements,'' into two items, one for securitization credit enhancements that are on-balance sheet assets and another for other credit enhancements. This will be accomplished by adding a new item 2.c., ''Standby letters of credit and other enhancements,'' where branches and agencies will disclose the unused portion of standby letters of credit and the maximum contractual amount of recourse or other credit exposure not in the form of an on-balance sheet asset that has been provided or retained in connection with the securitization structures reported in item 1 of Schedule S. This revision will enable the agencies to better understand the types of credit support that branches and agencies are providing to their securitizations, including which types are typically used for different types of securitized loans. The revisions will also achieve consistency with the changes to the Reports of Condition and Income (Call Report) filed by insured commercial banks and FDIC-supervised savings banks.

Request for Comment

Comments submitted in response to this Notice will be shared among the agencies and will be summarized or included in the Board's request for OMB approval. All comments will become a matter of public record. Written comments should address the accuracy of the burden estimates and ways to minimize burden as well as other relevant aspects of the information collection requests. Comments are invited on:

(a) Whether the proposed collection of information is necessary for the proper performance of the agencies' functions, including whether the information has practical utility;

(b) The accuracy of the agencies' estimate of the burden of the information collection, including the validity of the methodology and assumptions used;

(c) Ways to enhance the quality, utility, and clarity of the information to be collected;

(d) Ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology; and

(e) Estimates of capital or start up costs and costs of operation, maintenance, and purchase of services to provide information.

Board of Governors of the Federal Reserve System, February 27, 2003.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 03–5070 Filed 3–4–03; 8:45 am]

BILLING CODE 6210–01–P

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 Web site 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 March 19, 2003.

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

1. Harrodsburg First Financial Bancorp, Inc., Harrodsburg, Kentucky; to engage *de novo* through its subsidiary, First South Credit of Versailles, Inc., Versailles, Kentucky, in consumer finance activities, pursuant to § 225.28(b)(1) of Regulation Y.

Board of Governors of the Federal Reserve System, February 27, 2003.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 03–5071 Filed 3–4–03; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–03–48]

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 the CDC Reports Clearance Officer on (404) 498–1210.

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. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project

Performance Evaluation Program for Rapid HIV Testing—New—Public Health Practice Program Office (PHPPO), Centers for Disease Control and Prevention (CDC).

To support our mission of improving public health and preventing disease through continuously improving laboratory practices, the Model Performance Evaluation Program (MPEP), Division of Laboratory Systems, Public Health Practice Program Office, Centers for Disease Control and Prevention intends to provide a new HIV rapid testing performance evaluation program (HIV Rapid Testing MPEP). This program will offer external performance evaluation (PE) for rapid tests such as the OraQuick® Rapid HIV–1 Antibody Test, recently approved as a waived test by the U.S. Food and Drug Administration, and for other licensed tests such as the Abbott-Murex SUDS® HIV–1 Test. Participation in PE programs is expected to lead to improved HIV testing performance because participants have the opportunity to identify areas for improvement in testing practices. This program will help to ensure accurate testing as a basis for development of HIV prevention and intervention strategies.

This external quality assessment program will be made available at *no cost* (for receipt of sample panels) to sites performing rapid testing for HIV antibodies. This program will offer laboratories/testing sites an opportunity for:

(1) Assuring that the laboratories/testing sites are providing accurate tests through external quality assessment.

(2) Improving testing quality through self-evaluation in a non-regulatory environment.

(3) Testing well characterized samples from a source outside the test kit manufacturer.

(4) Discovering potential testing problems so that laboratories/testing sites can adjust procedures to eliminate them.

(5) Comparing individual laboratory/testing site results to others at a national and international level, and consulting with CDC staff to discuss testing issues.

Participants in the MPEP HIV Rapid Testing program will be required to complete a laboratory practices questionnaire survey annually. In addition, participants will be required to submit results twice/year after testing mailed performance evaluation samples. The estimated annualized cost to respondents is $2,625.00.

Forms	Number of respondents	Frequency of responses	Average burden/response (in hours)	Total burden (in hours)
HIV Rapid Testing Questionnaire	300	1	15/60	75
HIV Rapid Testing Results Booklet	300	2	10/60	100
Total				175

Dated: February 27, 2003.

Thomas Bartenfeld,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 03–5120 Filed 3–4–03; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Dermatologic and Ophthalmic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Dermatologic and Ophthalmic Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on March 17, 2003, from 8 a.m. to 5:30 p.m.

Location: Holiday Inn, The Ballrooms, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Kimberly Littleton Topper, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7001, e-mail: topperk@cder.fda.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12534. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss new drug application (NDA) 21–414, VITRASE (hyaluronidase for intravitreal injection), ISTA Pharmaceuticals, for the treatment of vitreous hemorrhage. The background material for this meeting will be posted on the Internet when available or 1 working day before the meeting at http://www.fda.ohrms/dockets/ac/menu.htm.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by March 10, 2003. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before March 10, 2003, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Kimberly Littleton Topper at least 7 days in advance of the meeting.

FDA regrets that it was unable to publish this notice 15 days prior to the March 17, 2003, Dermatologic and Ophthalmic Drugs Advisory Committee meeting. Because the agency believes there is some urgency to bring this issue to public discussion and qualified members of the Dermatologic and Ophthalmic Drugs Advisory Committee