Estimated Time per Response: 4 hours.
Frequency of Response: On occasion reporting requirement.
Obligation to Respond: Required to obtain or retain benefits.

Total Annual Burden: 1,120 hours.
Annual Cost Burden: $16,500.

Privacy Act Impact Assessment: N/A.

Nature and Extent of Confidentiality: Applicants may request that information be withheld from public inspection pursuant to 47 CFR 0.459 of the Commission’s rules. The request must be justified pursuant to 47 CFR 0.457.

Needs and Uses: This collection will be submitted as an extension (no change in reporting requirements) after this 60 day comment period to the Office of Management and Budget (OMB) in order to obtain the full three year clearance. Mandatory electronic filing of applications for Experimental Radio licenses, including FCC Form 442, commenced on January 1, 2004. Applicants that required an FCC license to operate a new or modified experimental radio station must file FCC Form 442, as required by 47 CFR 5.55(a)–(c) and 47 CFR 5.59 of the Commission’s rules. The FCC’s information technician and engineers use the data supplied by applicants in the FCC Form 442 to determine: (1) If the applicant is eligible for an experimental license; (2) the purpose of the experiment; (3) compliance with the requirements of Part 5 of the Commission’s rules; and (4) if the proposed operation will cause interference to existing operations. Thus, the FCC cannot grant an experimental license without the information contained on this form.

Marlene H. Dortch,
Secretary.
[FR Doc. E8–5764 Filed 3–19–08; 8:45 am]
BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Reviewed by the Federal Communications Commission for Extension Under Delegated Authority, Comments Requested

March 13, 2008.

SUMMARY: The Federal Communications Commission, as part of its continuing effort to reduce paperwork burdens, invites the general public and other Federal agencies to take this opportunity to (PRA) of 1995 (PRA), Public Law 104–13. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. Subject to the PRA, no person shall be subject to any penalty for failing to comply with a collection of information that does not display a valid control number.

Comments are requested concerning (a) whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission’s burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology.

DATES: Written PRA comments should be submitted on or before May 19, 2008. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: You may submit all PRA comments by e-mail or U.S. post mail.

To submit your comments by e-mail, send an e-mail to PRA@fcc.gov. To submit your comments by U.S. mail, mark them to the attention of Cathy Williams, Federal Communications Commission, Room 1–C323, 445 12th Street, SW., Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: For additional information about the collection(s), contact Cathy Williams at (202) 418–2918 or send an e-mail to PRA@fcc.gov.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0236.
Title: Section 74.703, Interference.
Form Number: Not applicable.
Type of Review: Extension of a currently approved collection.
Respondents: Business or other for-profit entities, Not-for-profit institutions, State, local or tribal government.
Number of Respondents: 100.
Estimated Time per Response: 10 hours.
Frequency of Response: On occasion reporting requirement.

Obligation to Respond: Required to obtain or retain benefits.

Confidentiality: No need for confidentiality required.
low power TV or TV translator operating on the same channel or first adjacent channel of its intention to initiate or change wireless operations and the likelihood of interference from the low power TV or translator station within its licensed geographic service area. The notice should describe the facilities, associated service area and operations of the wireless licensee with sufficient detail to permit an evaluation of the likelihood of interference. Upon receipt of such notice, the digital LPTV or TV translator licensee must cease operation within 120 days unless: (1) It obtains the agreement of the wireless licensee to continue operations; (2) the commencement or modification of wireless service is delayed beyond that period (in which case the period will be extended); or (3) the Commission stays the effect of the interference notification, upon request.

47 CFR 74.703(h) requires in each instance where suspension of operation is required, the licensee shall submit a full report to the FCC in Washington, DC, after operation is resumed, containing details of the nature of the interference, the source of the interfering signals, and the remedial steps taken to eliminate the interference.

Federal Communications Commission.

Marlene H. Dortch,
Secretary.

[FR Doc. E8–5770 Filed 3–19–08; 8:45 am]
BILLING CODE 6712–01–P

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 applications 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 April 14, 2008.

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

1. Select Bancorp, Inc.; to become a bank holding company by acquiring 100 percent of the voting shares of Select Bank & Trust Company, both of Greenville, North Carolina.

B. Federal Reserve Bank of St. Louis (Glenda Wilson, Community Affairs Officer) 411 Locust Street, St. Louis, Missouri 63101–2034:

1. Cross County Bancshares, Inc., Wynne, Arkansas; to acquire additional voting shares of First Southern Bank, Batesville, Arkansas, for a total of up to 13.13 percent.

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

1. CTB Financial Corporation, Ruston, Louisiana; to acquire 100 percent of the voting shares of Community Trust Bank of Texas, Dallas, Texas, a de novo bank.


Robert dev. Frierson,
Deputy Secretary of the Board.

[FR Doc. E8–5630 Filed 3–19–08; 8:45 am]
BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Vision Health: Developing an Integrative Approach to Promotion and Protection, Request for Application (RFA) DP08–001

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 aforementioned meeting.

Time and Date: 12:30 p.m.–3:30 p.m., April 17, 2008 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of “Vision Health: Developing an Integrative Approach to Promotion and Protection, RFA DP08–001."

Contact Person for More Information: Susan B. Stanton, D.D.S., Scientific Review Administrator, CDC, 1600 Clifton Road, NE, Mailstop D72, Atlanta, GA 30333, Telephone: (404) 639–4640.

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.


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

[FR Doc. E8–5628 Filed 3–19–08; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0120]

Standards for Standardized Numerical Identifier, Validation, Track and Trace, and Authentication for Prescription Drugs; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is seeking information and comments on issues related to standards for identification, validation, tracking and tracing, and authentication for prescription drug products. Particularly, we are requesting information and comments from drug manufacturers, distributors, pharmacies, other supply chain stakeholders, foreign regulators, standards organizations, and other Federal agencies and interested parties. This request is related to FDA’s implementation of the Food and Drug