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 July 16, 1998.

A. Federal Reserve Bank of Boston (Richard Walker, Community Affairs Officer) 600 Atlantic Avenue, Boston, Massachusetts 02106-2204:

1. Danvers Bancorp, Inc., Danvers, Massachusetts; to become a bank holding company by acquiring 100 percent of the voting shares of Danvers Savings Bank, Danvers, Massachusetts.

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

1. One Valley Bancorp, Inc., Charleston, West Virginia; to acquire 100 percent of the voting shares of Summit Bankshares, Inc., Raphine, Virginia, and thereby indirectly acquire Bank of Rockbridge, Raphine, Virginia.

C. Federal Reserve Bank of Chicago (Philip Jackson, Applications Officer) 230 South LaSalle Street, Chicago, Illinois 60690-1413:

1. First American Bankshares, Inc., Fort Atkinson, Wisconsin; to acquire 100 percent of the voting shares of Jefferson County Bancorp, Inc., Jefferson, Wisconsin, and thereby indirectly acquire Jefferson County Bank, Jefferson, Wisconsin.

Board of Governors of the Federal Reserve System, June 16, 1998.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–16423 Filed 6–19–98; 8:45 am]

BILLING CODE 6210–01–F

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. Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

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 July 17, 1998.

A. Federal Reserve Bank of Boston (Richard Walker, Community Affairs Officer) 600 Atlantic Avenue, Boston, Massachusetts 02106-2204:

1. UST Corp., Boston, Massachusetts; to acquire and thereby merge with Affiliated Community Bancorp, Waltham, Massachusetts, and thereby indirectly acquire Lexington Savings Bank, Lexington, Massachusetts; and Middlesex Bank & Trust Company, Newton, Massachusetts.

In connection with this application, Applicant also has filed to acquire the Federal Savings Bank, Waltham, Massachusetts, and thereby operate a federal savings bank, pursuant to § 225.28(b)(4) of Regulation Y.

B. Federal Reserve Bank of Cleveland (Paul Kaboth, Banking Supervisor) 1455 East Sixth Street, Cleveland, Ohio 44101-2566:

1. Second Bancorp Incorporated, Warren, Ohio; to merge with Enfin, Inc., Solon, Ohio, and thereby indirectly acquire Enterprise Bank, Solon, Ohio.

C. Federal Reserve Bank of Atlanta (Lois Berthaume, Vice President) 104 Marietta Street, N.W., Atlanta, Georgia 30303-2713:

1. First American Corporation, Nashville, Tennessee; to acquire 100 percent of the voting shares of The Middle Tennessee Bank, Columbia, Tennessee.

2. Synovus Financial Corp., and TB&C Bancshares, Inc., both of Columbus, Georgia; to merge with Community Bank Capital Corporation, Alpharetta, Georgia, and thereby indirectly acquire Bank of North Georgia, Alpharetta, Georgia.

D. Federal Reserve Bank of Chicago (Philip Jackson, Applications Officer) 230 South LaSalle Street, Chicago, Illinois 60690-1413:

1. The Connor Trusts, Marshfield, Wisconsin; to acquire 51 percent of the voting shares of Pioneer Bancorp, Inc., Auburndale, Wisconsin, and thereby indirectly acquire Pioneer Bank, Auburndale, Wisconsin.

E. Federal Reserve Bank of St. Louis (Randall C. Sumner, Vice President) 411 Locust Street, St. Louis, Missouri 63102-2034:

1. National City Bancshares, Inc., Evansville, Indiana; to merge with Hoosier Hills Financial Corporation, Osgood, Indiana, and thereby indirectly acquire The Ripley County Bank, Osgood, Indiana.

Board of Governors of the Federal Reserve System, June 17, 1998.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–16543 Filed 6–19–98; 8:45 am]

BILLING CODE 6210–01–F

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.

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 July 7, 1998.

A. Federal Reserve Bank of New York (Betsy Buttrill White, Senior Vice President) 33 Liberty Street, New York, New York 10045-0001:

1. Deutsche Bank AG, Frankfurt am Main, Germany; to acquire German American Capital Corporation, New York, New York, and thereby engage in acquiring debt that is in default at the time of acquisition, pursuant to § 225.28(b)(2)(vii) of Regulation Y.

Board of Governors of the Federal Reserve System, June 17, 1998.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–16541 Filed 6–19–98; 8:45 am]

BILLING CODE 6210–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0364]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed reinstatement of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on product specific reports and recordkeeping requirements for certain electronic products.

DATES: Submit written comments on the collection of information by August 21, 1998.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Reporting and Recordkeeping for Electronic Products: Specific Product Requirements 21 CFR Parts 1020, 1030, 1040, and 1050 (OMB Control Number 0910–0213—Reinstatement)

Under sections 532 to 542 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360ii to 360ss), FDA has the responsibility to protect the public from unnecessary exposure to radiation from electronic products. Section 532 of the act (21 U.S.C. 360ii) directs the Secretary of the Department of Health and Human Services (the Secretary) to establish and carry out an electronic product radiation control program designed to protect the public health and safety from electronic radiation by, among other things, developing and administering performance standards for electronic products. Section 534(g) of the act (21 U.S.C. 360kk(g)) directs the Secretary to review and evaluate industry testing programs on a continuing basis; and section 535(e) and (f) of the act (21 U.S.C. 360ll(e) and (f)) directs the Secretary to immediately notify manufacturers of, and assure correction of, radiation defects or noncompliance with performance standards. The agency's authority to require records and reports is contained in section 537(b) and (c) of the act (21 U.S.C. 360nn(b) and (c)).

Under this authority, FDA issued regulations detailing product-specific performance standards that specify information to be supplied with the product or require specific reports.

The information collections are either specifically called for in the act or were developed to aid the agency in performing its obligations under the act. The data reported to FDA and the records that are maintained are used by FDA and the industry to make decisions and take actions that protect the public from radiation hazards presented by electronic products. This information refers to the identification of, location of, operational characteristics of, quality assurance programs for, and problem identification and correction of electronic products. The data provided to users and others are intended to encourage actions to reduce or eliminate radiation exposures.

The consequence of not obtaining the required information is that the public unknowingly may be exposed to unnecessary radiation hazards presented by electronic products. Without this information, FDA could not adequately make rational decisions and take appropriate actions to protect the public from these hazards as called for in the act.

Respondents to this collection of information are manufacturers, importers, and assemblers of electronic products. Not all of the requirements are placed on all of these groups.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
1020.20(c)(4)	1	1	1	1	1
1020.30(g)	200	1.33	265	35	9,275