1. Fifth Third Bancorp, Cincinnati, Ohio; to acquire Emerald Financial Corp., Strongville, Ohio, and thereby indirectly acquire Strongville Savings Bank, Strongville, Ohio, and thereby engage in savings and loan activities, pursuant to § 225.28(b)(4)(ii) of Regulation Y.

Board of Governors of the Federal Reserve System, April 20, 1999.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 99–10271 Filed 4–22–99; 8:45 am]

BILLING CODE 6210–01–F

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 10 a.m., Wednesday, April 28, 1999.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, N.W., Washington, DC 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any matters carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board; 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board's Web site at http://www.federalreserve.gov for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: April 21, 1999.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 99–10339 Filed 4–21–99; 10:03 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–0803]

Agency Information Collection Activities: Proposed Collection; Comment Request; Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents

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 a notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the proposed collection of information concerning restrictions on the sale and distribution of cigarettes and smokeless tobacco in order to protect children and adolescents under the Federal Food, Drug, and Cosmetic Act (the act).

DATES: Submit written comments on the collection of information by June 22, 1999.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Heather M. Rubino, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 15–74, Rockville, MD 20857, 301–827–3322.

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 extension 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 set forth in this document.

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.

Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents—OMB No. 0910–0312—Extension

Part 897 (21 CFR part 897) reflects requirements in sections 502(e)(2) and 520(e) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 352(e)(2) and 360j(e)).

Section 897.24 is intended to implement section 502(e)(2) of the act. Under section 502(e)(2) of the act, a device is misbranded unless its label bears the product's established name. Section 502(e)(4) of the act, in turn, explains that the ''established name'' with respect to a device means: (1) The applicable official name of the device designated under section 508 of the act (21 U.S.C. 358), (2) if there is no such name and the device is recognized in an official compendium, then the official title in such compendium, or (3) if neither (1) nor (2) apply, then ''any common or usual name of such device.'' Here, no official names have been designated under section 508 of the act, and these products are not recognized in an official compendium. Consequently, FDA developed established names for these products under section 502(e)(4) of the act. Section 897.24 requires that each cigarette or smokeless tobacco product package, carton, box, or container of any kind that is offered for sale, sold, or otherwise distributed bear