—Proposed Rule—Distressed Loan Restructuring

Dated: December 18, 2002.

Jeanette C. Brinkley,

Acting Secretary, Farm Credit Administration Board.

[FR Doc. 02–32373 Filed 12–19–02; 11:19 am]

BILLING CODE 6705–01–P

FEDERAL RESERVE SYSTEM

Agency Information Collection Activities: Announcement of Board Approval Under Delegated Authority and Submission to OMB

AGENCY: Board of Governors of the Federal Reserve System.

SUMMARY: *Background*: Notice is hereby given of the final approval of proposed information collections by the Board of Governors of the Federal Reserve System (Board) under OMB delegated authority, as per 5 CFR 1320.16 (OMB Regulations on Controlling Paperwork Burdens on the Public). Board–approved collections of information are incorporated into the official OMB inventory of currently approved collections of information. Copies of the OMB 83–I's and supporting statements and approved collection of information instrument(s) are placed into OMB's public docket files. The Federal Reserve may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

FOR FURTHER INFORMATION CONTACT: Federal Reserve Board Clearance Officer –Cindy Ayouch—Division of Research and Statistics, Board of Governors of the Federal Reserve System, Mail stop 41, Washington, DC 20551 (202–452–3829). OMB Desk Officer–Joseph Lackey—Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Final approval under OMB delegated authority of the extension for three years, without revision, of the following reports:

1. Report title: Recordkeeping and Disclosure Requirements Associated with Loans Secured by Real Estate Located in Flood Hazard Areas Pursuant to Section 208.25 of Regulation H.

Agency form number: Reg H–2.

OMB Control number: 7100–0280.

Frequency: Event–generated.

Reporters: State member banks.

Annual reporting hours: 111,420 hours.

Estimated average hours per response: Notice of special flood hazards to borrowers and servicers, 0.08 hours; notice to the Federal Emergency Management Agency (FEMA) of servicer, 0.08 hours; notice to FEMA of change of servicer, 0.08 hours; and retention of standard FEMA form, 0.04 hours.

Number of respondents: 976.

Small businesses are affected.

General description of report: This information collection is mandatory (12 U.S.C. 248(a)(1)). Because the Federal Reserve does not collect any of FEMA forms this information collection is not given confidential treatment. However, should any of these records come into the possession of the Federal Reserve, such information may be protected from disclosure by exemptions 4 and 6 of the Freedom of Information Act (5 U.S.C. 552(b)(4) and (b)(6)).

Abstract: Regulation H requires state member banks to notify a borrower and servicer when loans secured by real estate are determined to be in a special flood hazard area and notify them whether flood insurance is available; notify FEMA of the identity of, and any change of, the servicer of a loan secured by real estate in a special flood hazard area; and retain a completed copy of the Standard Flood Hazard Determination Form used to determine whether property securing a loan is in a special flood hazard area.

Board of Governors of the Federal Reserve System, December 17, 2002.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 02–32185 Filed 12–20–02; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: Federal Parent Locator Service.

OMB No. 0970–0142.

Description: State and local child support enforcement agencies may request the Federal Parent Locator Service (FPLS) to assist in locating parents in order to establish or enforce child support. The FPLS serves as a conduit between child support enforcement offices and Federal and state agencies by conducting weekly, biweekly, or monthly matches of the collected information with various agencies and distributing the information back to the requesting state or local child support office.

Respondents: State and local IV–D child support offices.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
FPLS submissions	5	24	1	120
Estimated Total Annual Burden Hours				120

In compliance with the requirements of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments 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)

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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: December 16, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–32183 Filed 12–20–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0063]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Consumer Surveys on Food and Dietary Supplement Labeling Issues

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (PRA).

DATES: Submit written comments on the collection of information by January 22, 2003.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Stuart Shapiro, Desk Officer for FDA.

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

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Consumer Surveys on Food and Dietary Supplement Labeling Issues—(OMB Control Number 0910–0492)—Extension

FDA is requesting an extension of the OMB approval of consumer surveys to help FDA's Center for Food Safety and Applied Nutrition formulate decisions and policies affecting the labeling of conventional foods and dietary supplements. Determining how consumers are likely to interpret various kinds of claims, disclaimers, warnings, caution statements, and notice statements that might appear in labeling is critical to agency decisionmaking under the Federal Food, Drug, and Cosmetic Act and the first amendment. It is often necessary to test actual or proposed labeling statements in realistic situations with typical consumers to determine what these label statements are communicating to consumers.

FDA or its contractor will collect and use information gathered from telephone, mail, shopping mall intercept, or Internet surveys to evaluate how consumers understand and respond to existing label statements, label statements proposed by industry or consumers, and other label statements that are under consideration as part of FDA's policy development process. Potential respondents to the surveys will be individual consumers either randomly chosen to represent specified populations or randomly assigned to experimental treatment conditions to control for the effects of individual differences in the population on the interpretation of label statements. In all instances, FDA will strive to collect a representative sample of individuals from the overall population or from relevant population groups as appropriate. FDA's general selection method will use stratification, with random sampling within the strata, to achieve representativeness for both overall populations and sensitive subpopulations, such as at-risk individuals or user segments. In the rare cases where geography is a limiting factor, FDA will use population-based cluster sampling to limit Government expense while preserving the statistical properties of the sample.

Respondents will provide background information and respond to package labels that contain the variations of label statements to be tested. Measures will include both self-reported comprehension and acceptance, as well as direct behavioral measures of consumer use and understanding of the package labeling.

FDA will use the information from the surveys in evaluating regulatory and policy options with respect to labeling. The agency often lacks empirical data about how consumers understand and respond to statements they might see in product labeling. The information gathered from such surveys can be used to test consumer comprehension and behavioral impact of various label statements and formats, taking into account the existing distribution of behavior, knowledge, and attitudes in the population that provides the context for understanding such statements. The surveys will help FDA assess consumer reactions to existing and proposed label statements.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Type of Survey	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Mail questionnaire	1,000	1	1,000	1	1,000
Telephone survey	2,000	1	2,000	.5	1,000
Internet or mail intercept survey	4,000	1	4,000	.5	2,000
Total					4,000

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

These estimates assume that as many as one mail survey project, one telephone survey project, and two Internet or mail intercept survey projects may be done on an annual basis. Estimates are based on the expected number of respondents necessary to obtain a statistically significant representation of important