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 July 9, 2001.

A. Federal Reserve Bank of Minneapolis (JoAnne F. Lewellen, Assistant Vice President) 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291:

1. Citizens State Bancorporation, Grafton, North Dakota; to merge with Ideal Bancshares, Inc., West Fargo, North Dakota, and thereby indirectly acquire voting shares of First Capital Bank of North Dakota, West Fargo, North Dakota; Walhalla Bank Holding Company, Walhalla, North Dakota; First State Bank of Langdon, Langdon, North Dakota, and Walhalla State Bank, Walhalla, North Dakota.

Board of Governors of the Federal Reserve System, June 11, 2001.

Robert deV. Frierson

Associate Secretary of the Board.

[FR Doc. 01–15101 Filed 6–14–01; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

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

TIME AND DATE: 10:00 a.m., Wednesday, June 20, 2001.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, NW., 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 items carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Michelle A. Smith, 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: June 12, 2001.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 01–15274 Filed 6–13–01; 11:29 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request Proposed Projects:

Title: Computerized Support Enforcement Systems.

OMB No. 0980–0271.

Description: The information being collected is mandated by Section 454(16) which provides for the establishment and operation by the State agency, in accordance with an initial and annually updated advance planning document approved under section 452(d) of this state, of a statewide system meeting the requirements of section 454A. In addition, 454A(e)(1) requires that States create a State Case Registry (SCR) within their statewide automated child support systems, to include information on IV–D cases and non-IV–D orders established or modified in the State on or after October 1, 1998. Section 454A(e)(5) requires States to regularly update their cases in the SCR.

The data being collected for the Advance Planning Document is a combination of narrative, budget and schedules which are used to provide funding approvals on a annual basis and to monitor and oversee system development.

The data being collected for the State Case Registry is used to transmit mandatory data elements to the Federal Case Registry where it is used for matching against other data bases for the purposes of location of individuals, assets, employment and other child support related activities.

Respondents: The respondents are 54 State and Territorial Child Support Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
307.15 (APD)	2	1	240	480
307.15 (APDU)	54	1	60	3240
307.11(e)(1)(ii) Collection of non-IV–D data for SCR States	54	25,200	.046	62,597
307.11(e)(1)(ii) Collection of non-IV–D data for SCR-courts	3,045	447	.029	39,472
307.11(e)(3)(v) Collection of Child Data for IV–D cases for SRC-Courts	3,045	213	.083	53,833
307.11(f)(1) Case Data Transmitted from SCR to FCR: New cases and case updates	54	46,379	2.82	130,788
Total				290,410

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 Information Services, 370 L'Enfant Promenade, SW., Washington, D.C. 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: June 6, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–15123 Filed 6–14–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0249]

Agency Information Collection Activities; Proposed Collection; Comment Request; Consumer and Producer Surveys on Economic Issues

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 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 proposed voluntary surveys of consumers and producers in order to help FDA comply with Executive Order 12866, the Regulatory Flexibility Act (RFA), and the Small Business Regulatory Enforcement Fairness Act of 1996 (SBREFA).

DATES: Submit written or electronic comments on the collection of information by August 14, 2001.

ADDRESSES: Submit electronic comments on the collection of information to http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. 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: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1227.

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© 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.

Consumer and Producer Surveys on Economic Issues

Under section 903(d)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(d)(2)), FDA is authorized to conduct research relating to regulated articles and to collect information relating to responsibilities of the agency. Executive Order 12866, RFA, and SBREFA direct Federal agencies to conduct regulatory impact analysis, and to consider flexible regulatory approaches. In order to perform the mandatory analysis it is often necessary to survey: (1) Regulated producers to determine existing practices and the changes in those practices likely under various policy options, (2) both consumers and manufacturers to explore attitudes towards policy proposals, and (3) industry experts to solicit expert opinions. FDA is seeking OMB clearance to conduct future surveys to implement Executive Order 12866, RFA, and SBREFA. Participation in the surveys will be voluntary. This request covers regulated entities, such as food processors, dietary supplement manufacturers, health professionals or other experts, and consumers.

FDA will use the information gathered from these surveys to identify current business practices, expert opinion, and consumer or manufacturer attitudes towards existing or proposed policy. FDA projects approximately 2 to 6 surveys per year, with a sample of between 10 and 1,000 respondents each for mail and telephone surveys, and a sample of up to 3,000 respondents for cable or internet surveys.

FDA estimates the upper bound 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	Hours per Response	Total Hours
Mail questionnaire	1,000	1	3	3,000