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 December 13, 1996.

A. Federal Reserve Bank of Chicago (James A. Bluemle, Vice President) 230 South LaSalle Street, Chicago, Illinois 60690:

1. Heartland Financial USA, Inc., Dubuque, Iowa; to acquire Tri-State Community Credit Corporation, Dubuque, Iowa, and thereby engage in operating a consumer finance company, pursuant to § 225.25(b)(1)(i) of the Board's Regulation Y; and act as agent in the sale of insurance directly related to extensions of credit by the consumer finance company, pursuant to § 225.25(b)(8)(ii)(A-C) of the Board's Regulation Y.

Board of Governors of the Federal Reserve System, November 22, 1996.

William W. Wiles,

Secretary of the Board.

[FR Doc. 96–30435 Filed 11–27–96; 8:45 am]

BILLING CODE 6210–01–F

Sunshine Act Meeting

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

TIME AND DATE: 10:00 a.m., Wednesday, December 4, 1996.

PLACE: Marriner S. Eccles Federal Reserve Board Building, C Street entrance between 20th and 21st Streets, N.W., Washington, D.C. 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: Mr. Joseph R. Coyne, Assistant to the Board; (202) 452–3204. You may call (202) 452–3207, beginning at approximately 5 p.m. two business days before this meeting, for a recorded announcement of bank and bank holding company applications scheduled for the meeting.

Dated: November 26, 1996.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 96–30659 Filed 11–26–96; 2:54 pm]

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: Protection and Advocacy System Annual Statement of Objectives and Priorities.

Description: Section 142 (a)(2) of the DD Act requires the State Protection and Advocacy System (P&As) to develop, by January 1st of each year, a Statement of Objectives and Priorities (SOP) and provide an opportunity for the public to comment on it. The final statement must be submitted (along with the prior year's PPR) to the regional office of DHHS. The Statement will provide the public and the Department a better understanding of the operation of the advocacy services and provide more comprehensive reporting to Congress.

Respondents: State, Local or Tribal Govt.; individuals or households; and not-for-profit institutions.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
P&A SOP	55	1	40	2,200

Estimated Total Annual Burden Hours: 2,200.

Title: State Developmental Disabilities Council Three Year State Plan.

Description: Part B, Sections 122 and 124 of the DD Act requires that each State must prepare and submit to the Secretary, DHHS, and have in effect, a State Plan providing information on individuals with developmental disabilities within a particular State and a description of the service needs of individuals with developmental disabilities and their families. The plan sets forth the goals and specific objectives to be achieved by the State in meeting the service needs of this population. The Plan describes State priorities, strategies, and actions, and the allocation of funds to meet stated goals and objectives.

Respondents: State, Local or Tribal Govt.; individuals or housholds; and not-for-profit institutions.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Three Year State Plan	55	1	100	5,500

Estimated Total Annual Burden Hours: 5,500.

Title: Developmental Disabilities Annual Protection and Advocacy Program Performance Report.

Description: Section 107(b) of the DD Act requires that by January 1st of each year P&A system established in a State shall prepare and transmit to the Secretary a Report describing activities, accomplishments, and expenditures of the System during the preceding year. This Report will provide ADD with information needed to ascertain whether a State is fulfilling the requirements of Public Law 104–183.

The Report will provide ADD an overview of program trends and achievements and will enable ADD to respond to administration and congressional requests. It will also be used to submit an Annual Report to Congress.

Respondents: State, Local or Tribal Govt.; individuals or households; and not-for-profit institutions.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
DD P&A PPR	55	1	40	2,200

Estimated Total Annual Burden Hours: 2,200.

Title: State Developmental Disabilities Council Annual Program Performance Report.

Description: Section 107 of the DD Act requires the State DD Councils of each State to prepare and transmit to the Secretary, DHHS, an annual Report for the preceding fiscal year. It is to describe the activities and resultant accomplishments carried out with Part B funds received for the Federal fiscal year, and the general situation in the State for individuals with developmental disabilities. The information is necessary for annual technical assistance and monitoring, as well as preparation of the Secretary's Annual Report to the President, the Congress, and the National Council on Disabilities.

Respondents: State, Local or Tribal Govt.; individuals or households; and not-for-profit institutions.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
DD Council PPR	55	1	44	2,420

Estimated Total Annual Burden Hours: 2,420.

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, Division of Information Resource Management Services, 370 L'Enfant Promenade, S.W., 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: November 22, 1996.

Douglas J. Godesky,

Reports Clearance Officer.

[FR Doc. 96–30455 Filed 11–27–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96E–0196]

Determination of Regulatory Review Period for Purposes of Patent Extension; DOMITOR®

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for DOMITOR® and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Commissioner of Patents and Trademarks, Department of Commerce, for the extension of a patent which claims that animal drug product.

ADDRESSES: Written comments and petitions should be directed to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Brian J. Malkin, Office of Health Affairs (HFY–20), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–1382.

SUPPLEMENTARY INFORMATION: The Drug Price Competition and Patent Term Restoration Act of 1984 (Pub. L. 98–417) and the Generic Animal Drug and Patent Term Restoration Act (Pub. L. 100–670) generally provide that a patent may be extended for a period of up to 5 years so long as the patented item (human drug product, animal drug product, medical device, food additive, or color additive) was subject to regulatory review by FDA before the item was marketed. Under these acts, a product's regulatory review period forms the basis for determining the amount of extension an applicant may receive.