1. Simmons First National Corporation, Pine Bluff, Arkansas; to acquire 100 percent of the voting shares of First Bank of Arkansas, Russellville, Arkansas, and thereby indirectly acquire First Bank of Arkansas, Searcy, Arkansas.

D. Federal Reserve Bank of Minneapolis (Karen L. Grandstrand, Vice President) 250 Marquette Avenue, Minneapolis, Minnesota 55480-2171:

1. State Bank of Hawley Employee Stock Ownership Plan and Trust, Hawley, Minnesota; to acquire 32.8 percent of the voting shares of Bankshares of Hawley, Inc., Hawley, Minnesota, and thereby indirectly acquire State Bank of Hawley, Hawley, Minnesota.

E. Federal Reserve Bank of Dallas (Genie D. Short, Vice President) 2200 North Pearl Street, Dallas, Texas 75201-2272:

1. Moody Bancshares, Inc., Galveston, Texas, and Moody Bank Holding Company, Reno, Nevada; each to acquire an additional 0.38 percent, for a total of 25.4 percent, of the voting shares of The Moody National Bank of Galveston, Galveston, Texas.

2. New Woodson Bancshares, Inc., Graham, Texas; to become a bank holding company by acquiring 100 percent of the voting shares of Woodson Bancshares, Inc., Woodson, Texas, and thereby indirectly acquire First State Bancorp, Inc., Carson City, Nevada, and First State Bank, Graham, Texas.

Board of Governors of the Federal Reserve System, May 29, 1997.

William W. Wiles,

Secretary of the Board.

[FR Doc. 97–14476 Filed 6–3–97; 8:45 am]

BILLING CODE 6210–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Health Care Policy and Research

Advisory Committee Meeting

In accordance with section 10(a) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2) announcement is made of the following advisory committee scheduled to meet during the month of June 1997:

Name: Health Services Research Dissemination Study Section.

Date and Time: June 19, 1997, 7:30 a.m.

Place: Holiday Inn Bethesda, 8120 Wisconsin Avenue, Delaware Room, Bethesda, Maryland 20815.

Open June 19, 1997, 7:30 a.m. to 7:45 a.m.

Closed for remainder of meeting.

Purpose: The Study Section is charged with the review of and making recommendations on grant applications for Federal support of conferences, workshops, meetings, or projects related to dissemination and utilization of research findings.

Agenda: The open session of the meeting on June 19, from 7:30 a.m. to 7:45 a.m. will be devoted to a business meeting covering administrative matters. During the closed session, the panel will be reviewing and discussing grant applications dealing with health services research issues. In accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C., 552b(c)(6), the Administrator, Agency for Health Care Policy and Research, has made a formal determination that this latter session will be closed because the discussions are likely to reveal personal information concerning individuals associated with the grant applications. This information is exempt from mandatory disclosure.

Anyone wishing to obtain a roster of members, minutes of the meeting, or other relevant information should contact Carmen Johnson, Office of Scientific Affairs, Agency for Health Care Policy and Research, Suite 400, 2101 East Jefferson Street, Rockville, Maryland 20852, Telephone (301) 594–1449 x1613.

Agenda items for all meetings are subject to change as priorities dictate.

Dated: May 21, 1997.

John Eisenberg,

Administrator.

[FR Doc. 97–14545 Filed 6–3–97; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 DAY–11–97]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Office on (404) 639–7090. Send written comments to CDC, Desk Officer; Human Resources and Housing Branch, New Executive Office Building, Room 10235; Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project

1. Continuing Medical Education (CME) Activity Registration Form—(0923–0013)—Extension—The Agency for Toxic Substances and Disease Registry (ATSDR) is mandated pursuant to the 1980 Comprehensive Environmental Response Compensation and Liability Act (CERCLA) and its 1986 Amendments, The Superfund Amendments and Reauthorization Act (SARA), to prevent or mitigate adverse human health effects and diminished quality of life resulting from the exposure to hazardous substances into the environment. As stated in CERCLA, the Administrator of ATSDR is charged to ''assemble, develop as necessary, and distribute to the states, and upon request to medical colleges, physicians, and other health professionals, appropriate educational materials (including short courses) on this topic''.

The development and use of activity registration forms for documenting participation in these activities at these meetings is an integral part of this process. This attendance documentation process is required by the Accreditation Council for Continuing Medical Education (ACCME), the body that authorizes agencies and institutions to award nationally recognized continuing medical education (CME) credit. As a condition of relicensure, physicians in 40 states are required to participate in CME courses. Individual physicians in these states are required to submit the number of hours of CME credit to state boards of professional registration at the time of relicensure. Failure by the physician to provide this information in a timely fashion will result in suspension of professional licensure.

This request is for a 3-year extension of the current OMB approval of uniform CME activity registration forms—one machine entry form and the other manually entered—to serve as the initial step in the development of an attendance documentation system. The total annual burden hours are 83.

Respondents	Number of respondents	Number of responses/ respondent	Average burden/ response (in hrs.)	Total burden (in hrs.)
Manual Entry Registration Form	500	1	0.083	41.5
Scantron Registration Form	500	1	0.083	41.5

Dated: May 29, 1997.

Wilma G. Johnson,

Acting Associate Director for Policy Planning And Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–14531 Filed 6–3–97; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Cooperative Agreements for Community-Based Primary Prevention Programs to Prevent Intimate Partner Violence for a Safe America, Program Announcement 727: Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting.

Name: Disease, Disability, and Injury Prevention and Control SEP: Cooperative Agreements for Community-Based Primary Prevention Programs to Prevent Intimate Partner Violence for a Safe America, Program Announcement 727.

Time and Date: 8:30 a.m.–5 p.m., June 24–25, 1997.

Place: Ramada Plaza Hotel, 4001 Presidential Parkway, Atlanta, Georgia 30341.

Status: Closed.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement 727.

The meeting will be closed to the public in accordance with provisions set forth in section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463.

Contact Person for More Information: James S. Belloni, Associate Director, State and Community Activities, National Center for Injury Prevention and Control, CDC, M/S K02, 4770 Buford Highway, NE, Atlanta, Georgia 30341–3724, telephone 770/488–4538.

Dated: May 28, 1997.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–14524 Filed 6–3–97; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) Announces the Following Meeting

Name: Determination of Optimal Frequency and Minimum Power Requirements for a New Radio-Frequency-Controlled Electrical Injury Protection System study protocol peer review.

Time and Date: 8:30–11:30 a.m., June 24, 1997.

Location: Suncrest Facility, Large Conference Room, NIOSH, CDC, 3040 University Avenue, Morgantown, West Virginia 26505.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Purpose: Participants will provide NIOSH with their individual advice and comments regarding technical and scientific aspects of the NIOSH protocol Determination of Optimal Frequency and Minimum Power Requirements for a New Radio-Frequency-Controlled Electrical Injury Protection System. Peer review panelists will review the study protocol and provide individual advice on the conduct of the study. Viewpoints and suggestions from industry, labor, academia, other governmental agencies, and the public are invited.

CONTACT PERSON FOR ADDITIONAL INFORMATION: Shengke Zeng, NIOSH, CDC, M/S 119, 1095 Willowdale Road, Morgantown, West Virginia 26505, telephone (304) 285–5971.

Dated: May 29, 1997.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–14523 Filed 6–3–97; 8:45 am]

BILLING CODE 4160–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0180]

Agency Information Collection Activities: Proposed Collection; Comment Request; Reinstatement

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 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 requirements relating to the preapproved or emergency shipment of a blood product for manufacturing prior to completion of hepatitis B surface antigen (HBsAg) testing and shipment of a blood product for manufacturing when the donor is known to be reactive for HBsAg.

DATES: Submit written comments on the collection of information by August 4, 1997.

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. Wolff, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600