Ports Authority (''Respondent''). The complaint was served on January 20, 2000. Complainants allege that Respondent violated sections 10 (d)(1) and (d)(4) of the Shipping Act of 1984, 46 U.S.C. app. § § 1709 (d)(1) and (d)(4), and violated the terms of a Settlement Agreement in FMC Docket No. 95–10, by assessing excess dockage charges on the basis of a new vessel measurement system contrary to the terms of its tariffs, giving no notice of such changes, and not following procedures as set forth in the Settlement Agreement.

This proceeding has been assigned to the office of Administrative Law Judges. Hearing in this matter, if any is held, shall commence within the time limitations prescribed in 46 CFR 502.61, and only after consideration has been given by the parties and the presiding officer to the use of alternative forms of dispute resolution. The hearing shall include oral testimony and cross-examination in the discretion of the presiding officer only upon proper showing that there are genuine issues of material fact that cannot be resolved on the basis of sworn statements, affidavits, depositions, or other documents or that the nature of the matter in issue is such that an oral hearing and cross-examination are necessary for the development of an adequate record. Pursuant to the further terms of 46 CFR 502.61, the initial decision of the presiding officer in this proceeding shall be issued by January 22, 2001, and the final decision of the Commission shall be issued by May 22, 2001.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 00–1855 Filed 1–25–00; 8:45 am]

BILLING CODE 6730–01–M

FEDERAL MARITIME COMMISSION

Ocean Transportation Intermediary License Applicants

Notice is hereby given that the following applicants have filed with the Federal Maritime Commission applications for licenses as Non-Vessel Operating Common Carrier and Ocean Freight Forwarder—Ocean Transportation Intermediaries pursuant to section 19 of the Shipping Act of 1984 as amended (46 U.S.C. app. 1718 and 46 CFR 515).

Persons knowing of any reason why any of the following applicants should not receive a license are requested to contact the Office of Freight Forwarders, Federal Maritime Commission, Washington, D.C. 20573.

Non-Vessel-Operating Common Carrier Ocean Transportation Intermediary Applicants:

Senator International Ocean LLC, 11250 NW 25th Street, Suite 124, Miami, FL 33172. Officers: Christian M. Ollino, Vice President (Qualifying Individual), Mario Alfonso, President.

Cargo Management International, 401 N. Oak Street, Inglewood, CA 90302. Officer: Ray A. Vidal, C.E.O. (Qualifying Individual).

Non-Vessel-Operating Common Carrier and Ocean Freight Forwarder Transportation Intermediary Applicants:

Pioneer International Corp., 80 Everett Avenue, Suite 325, Chelsea, MA 02150. Officers: Pamela Ann Grzonka, Asst. Vice President (Qualifying Individual), David W. Maloney, President.

PMG Containerline, Inc., 6300 Hazeltine National Dr., Suite 100, Orlando, FL 32822. Officers: James G. Gain, President (Qualifying Individual), Thomas R. Murray, Vice President.

HR Services d/b/a HR Shipping Services, 211 North Union Street, Suite 100, Alexandria, VA 22314. Officer: Nigel J. McCallum, V.P. Operations (Qualifying Individual).

Ocean Freight Forwarders—Ocean Transportation Intermediary Applicants:

Senator International Freight Forwarding LLC, 11250 N.W. 25th Street, Suite 124, Miami, FL 33172. Officers: Mario Alfonso, President (Qualifying Individual), Uwe Kirschbaum, Chairman.

Commercial Transport Co., 16820 Lee Road, Humble, TX 77396, Robert R. Chapa, Sole Proprietor.

Martin E. Button, Inc., 55 New Montgomery Street, Suite 400, San Francisco, CA 94105. Officers: Jennifer Rixford, Secretary (Qualifying Individual), Martin E. Button, President.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 00–1857 Filed 1–25–00; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

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

TIME AND DATE: 12:00 noon, Monday, January 31, 2000.

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

STATUS: Closed.

MATTERS TO BE CONSIDERED: 1. Proposals concerning renovation of a Federal Reserve Bank building. (This item was originally announced for a closed meeting on January 24, 2000.)

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

3. 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: January 21, 2000.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 00–1891 Filed 1–21–00; 4:36 pm]

BILLING CODE 6210–01–P

GENERAL SERVICES ADMINISTRATION

Interagency Committee for Medical Records (ICMR); Cancellation of Medical Standard Form

AGENCY: General Services Administration.

ACTION: Notice.

SUMMARY: Because of low usage the following Standard Form is cancelled: SF 539, Medical Record—Abbreviated Medical Record.

Since the Department of Defense (DoD) is the only agency still using this form, they have created a new form—DD 2770, Abbreviated Medical Record. This form is available from the DoD's web page. Address: http://web1.whs.oad.mil/icdhome/DDEFORMS.HTM

DATE: Effective on January 26, 2000.

FOR FURTHER INFORMATION CONTACT: Ms. Barbara Williams, General Services Administration, (202) 501–0581.

Dated: January 11, 2000.

Barbara M. Williams,

Deputy Standard and Optional, Forms Management Officer.

[FR Doc. 00–1854 Filed 1–25–00; 8:45 am]

BILLING CODE 6820–34–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–00–20]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention is providing opportunity for public comment on proposed data collection projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639–7090.

Comments are invited 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) ways to enhance 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 for other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Projects

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. Other than their time, there will be no cost to the respondents.

Respondents	No. of respondents	No. of responses/ respondent	Avg. burden per response	Total burden
Manual Entry Registration Form	2,000	1	4/60	133
Scantron Registration Form	3,000	1	5/60	250
Total	.....	.....	.....	383

Dated: January 20, 2000.

Nancy Cheal,

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

[FR Doc. 00–1762 Filed 1–25–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0595]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Reporting and Recordkeeping Requirements for Manufacturers, Importers, User Facilities, and Distributors of Medical Devices Under FDAMA

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

DATES: Submit written comments on the collection of information by February 26, 2000.

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: Desk Officer for FDA.

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

SUPPLEMENTARY INFORMATION: In compliance with section 3507 of the PRA (44 U.S.C. 3507), FDA has