live via the Internet. To purchase these services call (703) 993–3100 or go to *http://www.capitolconnection.gmu.edu*.

Copies of materials adopted at this meeting can be purchased from the FCC's duplicating contractor, Best Copy and Printing, Inc. (202) 488–5300; Fax (202) 488–5563; TTY (202) 488–5562. These copies are available in paper format and alternative media, including large print/type; digital disk; and audio and video tape. Best Copy and Printing, Inc. may be reached by e-mail at *FCC@BCPIWEB.com*.

Marlene H. Dortch,

Secretary, Federal Communications Commission.

[FR Doc. E9–4888 Filed 3–4–09; 4:15 pm]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

Sunshine Act; Meeting

AGENCY HOLDING THE MEETING: Federal Maritime Commission.

TIME AND DATE: March 11, 2009–10 a.m.

PLACE: 800 North Capitol Street, NW., First Floor Hearing Room, Washington, DC.

STATUS: A portion of the meeting will be in Open Session and the remainder of the meeting will be in Closed Session.

MATTERS TO BE CONSIDERED:

Open Session

1. Docket No. 02–15 Passenger Vessel Financial Responsibility—Request of Commissioner Brennan.

2. Anderson International Transport and Owen Anderson, *et al.*—Order of Investigation and Hearing—Request for Extension of Time.

3. FY 2008 Federal Human Capital Survey Results—Interpretation of Results & Trends Closed Session.

4. Internal Administrative Practices and Personnel Matters.

CONTACT PERSON FOR MORE INFORMATION: Karen V. Gregory, Secretary, (202) 523–5725.

Karen V. Gregory,

Secretary.

[FR Doc. E9–4949 Filed 3–4–09; 4:15 pm]

BILLING CODE 6730–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10110, CMS–R–250, CMS–R–144/CMS–368 and CMS–668B]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the Agency's function; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Manufacturer Submission of Average Sales Price (ASP) Data for Medicare Part B Drugs and Biologicals; *Use:* Section 1847A of the Social Security Act requires that the Medicare Part B payment amounts for covered drugs and biologicals not paid on a cost or prospective payment basis be based upon manufacturers' average sales price data submitted to CMS. CMS will utilize the ASP data to determine the Medicare Part B drug payment amounts. *Form Number:* CMS–10110 (OMB# 0938–0921); *Frequency:* Quarterly; *Affected Public:* Business or other for-profits; *Number of Respondents:* 180; *Total Annual Responses:* 720; *Total Annual Hours:* 28,800. (For policy questions regarding this collection contact Catherine Jansto at 410–786–7762. For all other issues call 410–786–1326.)

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* MPAF Data and Supporting Regulations in 42 CFR 413.337, 413.343, 424.32 and 483.20; *Use:* Resident assessment information that Skilled Nursing Facilities (SNFs) are required to submit is described under section 42 CFR 413.343 and 483.20. The manner necessary to administer the payment rate methodology is described under section 42 CFR 413.337. An assessment form comprised of a subset of resident assessment information has been developed for use by SNFs to satisfy Medicare payment requirements, in lieu of a full Minimum Data Set. The associated burden is the time the SNF staff is required to complete the Medicare PPS Assessment Form (MPAF), SNF staff time to encode, and SNF staff time spent in transmitting the data. *Form Number:* CMS–R–250 (OMB# 0938–0739); *Frequency:* Occasionally; *Affected Public:* Business or other for-profits and not-for-profit institutions, State, Local, or Tribal Governments, and Federal Government; *Number of Respondents:* 15,039; *Total Annual Responses:* 3,834,945; *Total Annual Hours:* 2,704,764. (For policy questions regarding this collection contact Julie Stankivic at 410–786–5725. For all other issues call 410–786–1326.)

3. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* State Medicaid Drug Rebate; *Use:* Section 1927 of the Social Security Act requires each State Medicaid agency to report quarterly prescription drug utilization information to drug manufacturers and to CMS. As part of this information, the State Medicaid agencies are required to report the total Medicaid rebate amount they claim they are owed by each drug manufacturer for each covered prescription drug product each quarter. *Form Number:* CMS–R–144 and CMS–368 (OMB# 0938–0582); *Frequency:* Quarterly; *Affected Public:* State, Local or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 204; *Total Annual Hours:* 9,389. (For policy questions regarding this collection contact Dusty Kerhart at 410–786–3273. For all other issues call 410–786–1326.)

4. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Post Clinical Laboratory Survey Questionnaire and Supporting Regulations in 42 CFR 493.1771, 493.1773, and 493.1777; *Use:* This form is used by the State agency to determine a laboratory's compliance with the Clinical Laboratory Improvement Amendments of 1988 (CLIA). This information is needed for a laboratory's CLIA certification and recertification. *Form Number:* CMS–

668B (OMB# 0938–0653); *Frequency:* Biennially; *Affected Public:* Business or other for-profits and not-for-profit institutions. State, Local, or Tribal Government, Federal Government; *Number of Respondents:* 21,000; *Total Annual Responses:* 10,500; *Total Annual Hours:* 2,625. (For policy questions regarding this collection contact Kathleen Todd at 410–786–3385. For all other issues call 410–786–1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995*, or e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov*, or call the Reports Clearance Office on (410) 786–1326.

To be assured consideration, comments and recommendations for the proposed information collections must be received by the OMB desk officer at the address below, no later than 5 p.m. on *April 6, 2009*.

OMB, Office of Information and Regulatory Affairs, *Attention:* CMS Desk Officer, *Fax Number:* (202) 395–6974, E-mail: *OIRA_submission@omb.eop.gov*.

Dated: February 27, 2009.

Michelle Shortt,

Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. E9–4791 Filed 3–5–09; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10165]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Electronic Health Records (EHR) Demonstration Web Enabled Application for Phase II; *Use:* The goal of the Electronic Health Record (EHR) demonstration is to foster the implementation and adoption of EHRs and Health Information Technology (HIT) more broadly as effective vehicles improve the quality of care provided and transform the way medicine is practiced and delivered. Adoption of HIT has the potential to provide significant savings to the Medicare program and improve the quality of care rendered to Medicare beneficiaries. This demonstration is designed to leverage the combined forces of private and public payers to drive physician practices to widespread adoption and use of EHRs. The demonstration is being implemented in two phases. Over 800 applications were received, via a manual (paper) process, from interested practices in the four Phase I sites. Because of the greater number of sites and projected applicants for Phase II, CMS has Web enabled the application. This is expected to make it easier for practices to complete the application accurately and completely, submit it in a timely manner, and allow CMS to process the applications more efficiently and effectively. *Form Number:* CMS–10165(OMB#: 0938–0936–0965); *Frequency:* Reporting—Once; *Affected Public:* Business or other for-profit and not-for-profit institutions; *Number of Respondents:* 1,600; *Total Annual Responses:* 1,600; *Total Annual Hours:* 347. (For policy questions regarding this collection contact Jody Blatt at 410–786–6921. For all other issues call 410–786–1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS' Web Site at *http://www.cms.hhs.gov/PaperworkReductionActof1995*, or e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov*, or call the Reports Clearance Office on (410) 786–1326.

In commenting on the proposed information collections please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in one of the following ways by *May 5, 2009*:

1. *Electronically*. You may submit your comments electronically to *http://www.regulations.gov*. Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) accepting comments.

2. *By Regular Mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: February 27, 2009.

Michelle Shortt,

Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. E9–4807 Filed 3–5–09; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0097]

Agency Information Collection Activities; Proposed Collection; Comment Request; Request for Samples and Protocols

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 the information collection requirements relating to the regulations which state that protocols for samples of biological products must be submitted to the agency.

DATES: Submit written or electronic comments on the collection of information by May 5, 2009.