Sandra L. Kusumoto,

Director, Bureau of Certification and Licensing.

[FR Doc. 2010–7890 Filed 4–6–10; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL MARITIME COMMISSION

Ocean Transportation Intermediary License Revocation

The Federal Maritime Commission hereby gives notice that the following Ocean Transportation Intermediary licenses have been revoked pursuant to section 19 of the Shipping Act of 1984 (46 U.S.C. Chapter 409) and the regulations of the Commission pertaining to the licensing of Ocean Transportation Intermediaries, 46 CFR part 515, effective on the corresponding date shown below:

License Number: 021331F.

Name: Deseret Forwarding International, Inc.

Address: 1760 Airway, Suite 103, El Paso, TX 79925.

Date Revoked: February 25, 2010.

Reason: Failed to maintain a Valid Bond.

License Number: 004553N.

Name: Marianas Steamship Agencies, Inc. DBA MSA Logistics.

Address: Commercial Port Annex, 2nd Floor, 1010 Cabras Highway, Piti, Guam 96915.

Date Revoked: March 4, 2010.

Reason: Surrendered license voluntarily.

Sandra L. Kusumoto,

Director, Bureau of Certification and Licensing.

[FR Doc. 2010–7888 Filed 4–6–10; 8:45 am]

BILLING CODE 6730–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology HIT Policy Committee Advisory Meeting; Notice of Meeting

AGENCY: Office of the National Coordinator for Health Information Technology, HHS.

ACTION: Notice of change of location for meetings.

This notice references forthcoming meetings of public advisory committees of the Office of the National Coordinator for Health Information Technology (ONC). The meeting will be open to the public.

Name of Committee: HIT Policy Committee; Meaningful Use Workgroup.

General Function of the Committee: to provide recommendations to the National Coordinator on a policy framework for the development and adoption of a nationwide health information technology infrastructure that permits the electronic exchange and use of health information as is consistent with the Federal Health IT Strategic Plan and that includes recommendations on the areas in which standards, implementation specifications, and certification criteria are needed.

Date and Time: The meetings will be held on April 20, 2010, from 9 a.m. to 5 p.m./Eastern Time (the Meaningful Use Workgroup); and April 21, 2010, from 10 a.m. to 4 p.m./Eastern Time (HIT Policy Committee).

Location: The location for both meetings has changed to the Renaissance Dupont Circle Hotel, 1143 New Hampshire Avenue, NW., Washington, DC. The hotel telephone number is 202–775–0800.

Contact Person: Judy Sparrow, Office of the National Coordinator, HHS, 330 C Street, SW., Washington, DC 20201, 202–205–4528, Fax: 202–690–6079, e-mail: *judy.sparrow@hhs.gov* Please call the contact person for up-to-date information on these meetings. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice.

Agenda: The Meaningful Use Workgroup meeting will concern Patient/Consumer Engagement, and hear testimony from experts; the HIT Policy Committee will hear reports from its workgroups, including the Meaningful Use Workgroup, the Certification/Adoption Workgroup, the NHIN Workgroup, the Privacy & Security Policy Workgroup, and the Strategic Plan Workgroup. ONC intends to make background material available to the public no later than two (2) business days prior to the meeting. If ONC is unable to post the background material on its Web site prior to the meeting, it will be made publicly available at the location of the advisory committee meeting, and the background material will be posed on ONC's Web site after the meeting, at *http://healthit.hhs.gov*.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before April 13, 2010. Oral comments from the public will be scheduled between approximately 3:30 p.m. to 4 p.m. Time allotted for each presentation is limited to three minutes. If the number of speakers requesting to comment is greater than can be reasonably accommodated during the scheduled open public hearing session, ONC will take written comments after the meeting until close of business.

Persons attending ONC's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

ONC welcomes the attendance of the public at its advisory committee meetings. Seating is limited at the location, and ONC will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Judy Sparrow at least seven (7) days in advance of the meeting.

ONC is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://healthit.hhs.gov* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (Pub. L. 92–463, 5 U.S.C., App. 2).

Dated: April 1, 2010.

Judith Sparrow,

Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. 2010–7902 Filed 4–6–10; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; the Jackson Heart Study (JHS)

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval the information collection listed below. This proposed information collection was previously published in the **Federal Register** on January 13, 2010, page 1789, and allowed 60 days for public comment. No comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented