Number of Respondents and Responses: 50 respondents; 50 responses.

*Estimated Time per Response:* 20 minutes.

*Frequency of Response:* On occasion reporting requirement.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection is Section 154(i) of the Communications Act of 1934, as amended.

Total Annual Burden: 17 hours. Total Annual Cost: \$63,750. Privacy Act Impact Assessment: No impact(s).

Nature and Extent of Confidentiality: There is no need for confidentiality with this collection of information.

Needs and Uses: 47 CFR 73.3588 states whenever a petition to deny or an informal objection has been filed against any applications for renewal, new construction permits, modifications, and transfers/assignments, and the filing party seeks to dismiss or withdraw the petition to deny or the informal objection, either unilaterally or in exchange for financial consideration, that party must file with the Commission a request for approval of the dismissal or withdrawal. This request must include the following documents: (1) A copy of any written agreement related to the dismissal or withdrawal, (2) an affidavit stating that the petitioner has not received any consideration in excess of legitimate and prudent expenses in exchange for dismissing/withdrawing its petition, (3) an itemization of the expenses for which it is seeking reimbursement, and (4) the terms of any oral agreements related to the dismissal or withdrawal of the petitions to deny. Each remaining party to any written or oral agreement must submit an affidavit within 5 days of petitioner's request for approval stating that it has paid no consideration to the petitioner in excess of the petitioner's legitimate and prudent expenses. The affidavit must also include the terms of any oral agreements relating to the dismissal or withdrawal of the petition to denv.

OMB Control No.: 3060–0626. Title: Section 90.483, Permissible Methods and Requirements of Interconnecting Private and Public Systems of Communications.

Form No.: N/A.

*Type of Review:* Extension of a currently approved collection. *Respondents:* Business of other for-

profit entities. Number of Respondents and Responses: 100 respondents; 100

*Responses:* 100 respondents; 100 responses.

*Estimated Time per Response:* 1 hour. *Frequency of Response:* On occasion reporting requirements; Third party disclosure requirement.

*Obligation To Respond:* Required to obtain or retain benefits. The statutory authority for this collection of information is contained in Sections 4(i), 11, 303(g), 303(r), and 332(c)(7) of the Communications Act of 1934, as amended, 47 U.S.C. 154(i), 161, 303(g), 303(r), 332(c)(7).

Total Annual Burden: 100 hours. Annual Cost Burden: None. Privacy Act Impact Assessment: No impact(s).

*Nature and Extent of Confidentiality:* There is no need for confidentiality with this collection.

*Needs and Uses:* When a frequency is shared by more than one system, automatic monitoring equipment must be installed at the base station to prevent activation of the transmitter when signals of co-channel stations are present and activation would interfere with communications in progress. Licensees may operate without the monitoring equipment if they have obtained the consent of all co-channel licensees located within a 120 kilometer (75 mile) radius of the interconnected base station transmitter. A statement must be submitted to the Commission indicating that all co-channel licensees have consented to operate without the monitoring equipment. This information is necessary to ensure that licensees comply with the Commission's technical and operational rules, and to prevent activation of the transmitter when signals of co-channel stations are present and could possibly interfere with communications in process.

Federal Communications Commission.

## Marlene H. Dortch,

Secretary, Office of Secretary. [FR Doc. 2016–22194 Filed 9–14–16; 8:45 am] BILLING CODE 6712–01–P

### FEDERAL MARITIME COMMISSION

## **Sunshine Act Meeting**

**AGENCY HOLDING THE MEETING:** Federal Maritime Commission.

**TIME AND DATE:** September 20, 2016; 10:00 a.m.

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

**STATUS:** The first portion of the meeting will be held in Open Session; the second in Closed Session.

### MATTERS TO BE CONSIDERED:

## **Open Session**

- 1. Briefing by Commissioner Maffei on U.S./Japan Bilateral Discussions
- 2. Staff Briefing on Foreign-based NVOCC Registration Renewal Process (Form FMC–65)

## **Closed Session**

- 1. Staff Briefing on Hanjin Shipping Bankruptcy and Shipping Disruptions
- 2. Staff Briefing on the Maersk/MSC Vessel Sharing Agreement, FMC Agreement No. 012293

## CONTACT PERSON FOR MORE INFORMATION:

Rachel E. Dickon, Assistant Secretary, (202) 523–5725.

### Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2016–22299 Filed 9–13–16; 4:15 pm] BILLING CODE 6730–AA–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Disease Control and Prevention

[CDC-2016-0090, Docket Number NIOSH 288-A]

## A Performance Test Protocol for Closed System Transfer Devices Used During Pharmacy Compounding and Administration of Hazardous Drugs

**AGENCY:** National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice of public meeting and request for public comment on a draft testing protocol.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) announces a public meeting concerning a universal closed system drug-transfer device (CSTD) testing protocol entitled, *A Performance Test Protocol for Closed System Transfer Devices Used During Pharmacy Compounding and Administration of Hazardous Drugs, http://www.cdc.gov/niosh/topics/ hazdrug/default.html/.* 

This is an opportunity for public comment on the protocol, the proposed list of surrogates, and to respond to NIOSH questions regarding the protocol.

To view the protocol and related materials, visit *www.regulations.gov* and enter CDC–2016–0090 in the search field and click "Search."