Synopsis: The agreement would authorize the parties to charter space from one another in the trade between the U.S. and Europe, the Middle East and Asia.

Agreement No.: 012382.

Title: Crowley/King Ocean Space Charter Agreement.

Parties: Crowley Caribbean Services, LLC and King Ocean Services Limited, Inc.

Filing Party: Wayne R. Rohde, Esq.; Cozen O'Connor; 1200 19th Street NW., Washington, DC 20036.

Synopsis: The agreement would authorize King Ocean to charter space to Crowley in the trade between the U.S. East Coast on the one hand and Aruba, Bonaire and Curacao on the other hand.

By Order of the Federal Maritime Commission.

Dated: December 29, 2015.

Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2015–33083 Filed 1–4–16; 8:45 am]

BILLING CODE 6731–AA–P

FEDERAL MINE SAFETY AND HEALTH REVIEW COMMISSION

Sunshine Act Meeting Notice

December 30, 2015.

TIME AND DATE: 10:00 a.m., Wednesday, January 13, 2016.

PLACE: The Richard V. Backley Hearing Room, Room 511N, 1331 Pennsylvania Avenue NW., Washington, DC 20004 (enter from F Street entrance).

STATUS: Open.

MATTERS TO BE CONSIDERED: The Commission will hear oral argument in the matter *Secretary of Labor* v. *Hibbing Taconite Company,* Docket Nos. LAKE 2013–231–RM, *et al.* (Issues include whether the Judge erred in upholding failure to abate orders.)

Any person attending this oral argument who requires special accessibility features and/or auxiliary aids, such as sign language interpreters, must inform the Commission in advance of those needs. Subject to 29 CFR 2706.150(a)(3) and 2706.160(d).

CONTACT PERSON FOR MORE INFO: Emogene Johnson (202) 434–9935/(202) 708–9300 for TDD Relay/1–800–877–8339 for toll free.

Sarah L. Stewart,

Deputy General Counsel.

[FR Doc. 2015–33203 Filed 12–31–15; 11:15 am]

BILLING CODE 6735–01–P

FEDERAL MINE SAFETY AND HEALTH REVIEW COMMISSION

Sunshine Act Meeting Notice

December 30, 2015.

TIME AND DATE: 10:00 a.m., Thursday, January 14, 2016

PLACE: The Richard V. Backley Hearing Room, Room 511N, 1331 Pennsylvania Avenue NW., Washington, DC 20004 (enter from F Street entrance).

STATUS: Open.

MATTERS TO BE CONSIDERED: The Commission will consider and act upon the following in open session: *Secretary of Labor* v. *Hibbing Taconite Company,* Docket Nos. LAKE 2013–231–RM, et al. (Issues include whether the Judge erred in upholding failure to abate orders.)

Any person attending this meeting who requires special accessibility features and/or auxiliary aids, such as sign language interpreters, must inform the Commission in advance of those needs. Subject to 29 CFR 2706.150(a)(3) and 2706.160(d).

CONTACT PERSON FOR MORE INFO: Emogene Johnson (202) 434–9935/(202) 708–9300 for TDD Relay/1–800–877–8339 for toll free.

Sarah L. Stewart,

Deputy General Counsel.

[FR Doc. 2015–33201 Filed 12–31–15; 11:15 am]

BILLING CODE 6735–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2007–D–0256 (Formerly 2007D–0089)]

Agency Information Collection Activities: Proposed Collection; Comment Request; Draft Guidance for Industry and Review Staff on Target Product Profile—A Strategic Development Process Tool

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on the reporting requirements contained in the draft guidance for industry and review staff entitled ''Target Product Profile—A Strategic Development Process Tool.''

DATES: Submit either electronic or written comments on the collection of information by March 7, 2016.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- Federal eRulemaking Portal: *http://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *http://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *http://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- Mail/Hand delivery/Courier (for written/paper submissions): Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2007–D–0256 (formerly 2007D–0089) for ''Agency Information Collection Activities: Proposed Collection; Comment Request; Draft Guidance for Industry and Review Staff on Target Product Profile—A Strategic Development Process Tool.'' Received