ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email to *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418–2918.

SUPPLEMENTARY INFORMATION:

OMB Control No.: 3060–0865.

Title: Wireless Telecommunications Bureau Universal Licensing System Recordkeeping and Third Party Disclosure Requirements.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for-profit entities, Individuals or households, Not-for-profit institutions, and State, Local or Tribal Government.

Number of Respondents and Responses: 84,048 respondents; 84,050 responses.

Estimated Time per Response: .166 hours (10 minutes)—4 hours.

Frequency of Response: Recordkeeping and third-party disclosure requirements; on occasion reporting requirement.

Obligation To Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in 47 U.S.C. 154(i) and 309(j).

Total Annual Burden: 116,306 hours.

Annual Cost Burden: No cost.

Needs and Uses: The Commission will submit this information collection to the Office of Management and Budget (OMB) as an extension after this 60-day comment period to obtain the full three-year clearance from them.

The purpose of this information collection is to continually streamline and simplify processes for wireless applicants and licensees, who previously used a myriad of forms for various wireless services and types of requests, in order to provide the Commission information that has been collected in separate databases, each for a different group of services. Such processes have resulted in unreliable reporting, duplicate filings for the same licensees/applicants, and higher cost burdens to licensees/applicants. By streamlining the Universal Licensing System (ULS), the Commission eliminates the filing of duplicative applications for wireless carriers; increases the accuracy and reliability of licensing information; and enables all wireless applicants and licensees to file all licensing-related applications and other filings electronically, thus increasing the speed and efficiency of the application process. The ULS also benefits wireless applicants/licensees by reducing the cost of preparing applications, and speeds up the licensing process in that the Commission can introduce new entrants more quickly into this already competitive industry. Finally, ULS enhances the availability of licensing information to the public, which has access to all publicly available wireless licensing information on-line, including maps depicting a licensee's geographic service area.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2023–04595 Filed 3–6–23; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

[Docket No. 22–35]

M.E. Dey & Co., Inc. Complainant v. Hapag-Lloyd AG and Hapag-Lloyd (America) LLC, Respondents; Notice of Filing of Amended Complaint and Assignment

Served: March 1, 2023

Notice is given that an Amended Complaint has been filed with the Federal Maritime Commission (Commission) by M.E. Dey & Co., Inc. hereinafter ''Complainant,'' against Hapag-Lloyd AG and Hapag-Lloyd (America) LLC. (hereinafter ''Respondents''). Complainant states that it is organized and existing under the laws of Wisconsin a non-vessel-operating common carrier with a principal place of business in Milwaukee, Wisconsin. Complainant identifies the Hapag-Lloyd AG is a global ocean carrier with headquarters in Hamburg, Germany, and Hapag-Lloyd (America) LLC as a United States subsidiary and agent of Hapag AG located in Atlanta, Georgia.

Complainant alleges that Respondents violated 46 U.S.C. 41102(c) and 41104(a)(14) regarding their practices and the billing and payment of charges on the shipments of cargo, including demurrage and rail storage charges and the failure to provide chassis. An answer to the complaint is due to be filed with the Commission within twenty-five (25) days after the date of service. The full text of the complaint can be found in the Commission's Electronic Reading Room at *https://www2.fmc.gov/readingroom/proceeding/22-35/.*

This proceeding has been assigned to Office of Administrative Law Judges. The initial decision of the presiding officer in this proceeding shall be issued by December 26, 2023, and the final decision of the Commission shall be issued by July 10, 2024.

William Cody, Secretary.

[FR Doc. 2023–04596 Filed 3–6–23; 8:45 am]

BILLING CODE 6730–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–0008]

Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public.

DATES: The meeting will take place virtually on April 20, 2023, from 9 a.m. to 6 p.m. Eastern Time.

ADDRESSES: Please note that due to the impact of this COVID–19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. Answers to commonly asked questions, including information regarding special accommodations due to a disability, may be accessed at: *https://www.fda.gov/advisory-committees/about-advisory-committees/common-questions-and-answers-about-fda-advisory-committee-meetings.*

FOR FURTHER INFORMATION CONTACT: Akinola Awojope, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5214, Silver Spring, MD 20993–0002, *Akinola.Awojope@fda.hhs.gov,* 301–636–0512, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). 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. Therefore, you should always check the Agency's website at *https://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the