The relevant rules for purposes of this revised information collection are §§ 25.146(a)(3) and 25.289(a)(2).

The new information collection requirements in this collection remove previously required information collections that are no longer considered necessary. In addition, the new information is required to ensure that applicants are meeting the Commission's rules when they modify their systems without prior approval from the Commission. In addition, the information is required to ensure that the public and the Commission are provided with accurate information about the operations of a system. amended the registration and verification requirements for individual users.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2026–23042 Filed 11–10–26; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

[Docket No. 26–04]

Orleans International, Inc., Complainant v. Hapag Lloyd AG, Respondent; Notice of Filing of Second Amended Complaint

Notice is given that a second amended complaint (the ''amended complaint'') has been filed with the Federal Maritime Commission (the ''Commission'') by Orleans International, Inc. (the ''Complainant'') against Hapag Lloyd AG (the ''Respondent''). Complainant states that the Commission has subject-matter jurisdiction over the amended complaint pursuant to the Shipping Act of 1984, as amended, 46 U.S.C. 41301 *et seq.* and 41310, and personal jurisdiction over Respondent as a vessel-operating ocean common carrier, as defined in 46 U.S.C. 40102(18).

Complainant is a corporation existing under the laws of the state of Michigan with its principal place of business located in Farmington Hills, Michigan.

Complainant identifies Respondent Hapag Lloyd AG as a company existing under the laws of the Federal Republic of Germany with its principal place of business located in Hamburg, Germany, whose agent in the United States is Hapag-Lloyd (America) LLC, a limited liability company existing under the laws of the state of Delaware with its principal place of business located in Atlanta, Georgia.

Complainant alleges that Respondents violated 46 U.S.C. 41102(c); 41104(a)(10) and (14); and 46 CFR 545.5. Complainant alleges these violations arose from the assessment of demurrage and detention charges during periods of time in which Complainant's ability to pick up or return containers was constrained due to circumstances beyond its control, Respondent's lack of engagement in attempted dispute resolution, and other acts or omissions of Respondent.

An answer to the amended complaint must be filed with the Commission within 25 days after the date of service.

The full text of the amended complaint can be found in the Commission's electronic Reading Room at *https://www2.fmc.gov/readingroom/proceeding/26-04/.* This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding judge shall be issued by March 19, 2027, and the final decision of the Commission shall be issued by October 4, 2027.

(Authority: 46 U.S.C. 41301; 46 CFR 502.61(c))

Served: November 9, 2026.

Jennifer Everling,

Assistant Secretary.

[FR Doc. 2026–23065 Filed 11–10–26; 8:45 am]

BILLING CODE 6730–02–P

FEDERAL TRADE COMMISSION

SES Performance Review Board

AGENCY: Federal Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given of the appointment of members to the FTC Performance Review Board.

FOR FURTHER INFORMATION CONTACT: Tamika Williams, Chief Human Capital Officer, 600 Pennsylvania Avenue NW, Washington, DC 20580, (202) 326–2184.

SUPPLEMENTARY INFORMATION: Pursuant to 5 U.S.C. 4314(c)(4) and 5 CFR 430.311(a)(4), agencies must publish notice of Performance Review Board (PRB) appointments in the **Federal Register** before service begins. The PRB reviews and evaluates the initial appraisal of a senior executive's performance by the supervisor, and makes recommendations regarding performance ratings, performance-based pay adjustments, and awards (bonuses) to the Chairman.

The following individuals have been designated to serve on the Commission's Performance Review Board:

Lucas Croslow, General Counsel, PRB Chair

Daniel Guarnera, Director, Bureau of Competition

Christopher Mufarrige, Director, Bureau of Consumer Protection

Ted Rosenbaum, Deputy Director for Research and Management, Bureau of Economics

Brian Berggren, Associate Director for Enforcement, Bureau of Consumer Protection

Katherine White, Deputy Director, Bureau of Consumer Protection

Tamika Williams, Chief Human Capital Officer

Montreal Ravenel, Deputy Chief Human Capital Officer and Performance Management Officer

By direction of the Commission.

April J. Tabor,

Secretary.

[FR Doc. 2026–23028 Filed 11–10–26; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Final Effect of the Designation of a Class of Employees for Addition to the Special Exposure Cohort

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

ACTION: Notice.

SUMMARY: HHS gives notice of the final effect of a decision to add a class of employees from the Savannah River Site in Aiken, South Carolina, to the Special Exposure Cohort (SEC) under the Energy Employees Occupational Illness Compensation Program Act of 2000.

FOR FURTHER INFORMATION CONTACT: Lori Marion-Moss, Director, Division of Compensation Analysis and Support, NIOSH, 1090 Tusculum Avenue, MS C–46, Cincinnati, OH 45226–1938, Telephone 1–877–222–7570. Information requests can also be submitted by email to *DCAS@CDC.GOV.*

SUPPLEMENTARY INFORMATION:

Authority: 42 U.S.C. 7384q(b). 42 U.S.C. 7384l(14)(C).

On July 8, 2026, as provided for under 42 U.S.C. 7384l(14)(C), the Secretary of HHS designated the following class of employees as an addition to the SEC:

All construction trade employees of Department of Energy (DOE) subcontractors who worked at the Savannah River Site in Aiken, South Carolina, from January 1, 1991, through December 31, 1993, for a number of workdays aggregating at least 250 workdays, occurring either solely under this employment, or in combination with

workdays within the parameters established for one or more other classes of employees included in the Special Exposure Cohort. This class specifically does not include employees of prime contractors of Westinghouse Savannah River Company during this time period.

This designation became effective on August 7, 2026. Therefore, beginning on August 7, 2026, members of this class of employees, as defined in this notice, became members of the SEC.

John J. Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Department of Health and Human Services.

[FR Doc. 2026–23004 Filed 11–10–26; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10856]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by January 11, 2027.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send 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) that are 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.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request*: Revision of a currently approved collection; *Title of Information Collection*: Medicaid Managed Care and Supporting Regulations; *Use*: The collection of information request pertains to the attestation collection requirement at 42 CFR 438.6(c)(2)(ii)(H), which requires that providers receiving payment under a State directed payment attest that they do not participate in any hold harmless arrangement for any health care-related tax as specified in § 433.68(f)(3) in which the State or other unit of government imposing the tax provides for any direct or indirect payment, offset, or waiver such that the provision of the payment, offset, or waiver directly or indirectly guarantees to hold the taxpayer harmless for all or any portion of the tax amount, and ensure either that (upon CMS request) such attestations are available, or that the State provides an explanation that is satisfactory to CMS about why specific providers are unable or unwilling to make such attestations. *Form Number*: CMS–10856 (OMB control number: 0938–1453); *Frequency*: Yearly and once; *Affected Public*: Private sector and State, Local, or Tribal Governments; *Number of Respondents*: 719; *Total Annual Responses*: 6,243; *Total Annual Hours*: 264,468. (For questions regarding this collection contact Alex Loizias at 410–786–2435.)

William N. Parham, III

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2026–23037 Filed 11–10–26; 8:45 am]

BILLING CODE 4169–69–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–D–13432]

Early-Phase IND Protocol Recommendations for Solid Tumors; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for industry titled ''Early-Phase IND Protocol Recommendations for Solid Tumors.'' The purpose of this draft guidance is to provide recommendations to sponsors and clinical investigators developing early-phase trials, which include first-in-human (FIH) protocols or protocols for