wholesalers, flower, nursery stock, and florist's supplies wholesalers, state government, other chemical and allied product merchant wholesalers, exterminating and pest control service, management, scientific, and technical consulting services. North American Industrial Classification System (NAICS) codes identified in question 12 of the ICR.

Respondent's obligation to respond: Mandatory. FIFRA and FFDCA.

Estimated number of potential respondents: 15,023.

Frequency of response: On occasion.

Total estimated average number of responses for each respondent: Between 1 to 74.

Total estimated burden: 2,234,433 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated costs: $172,873,497 (per year), includes $0 annualized capital investment or maintenance and operational costs.

III. Are there changes in the estimates from the last approval?

There is an increase of 59,385 hours in the total estimated respondent burden compared with that identified in the ICR currently approved by OMB. This increase reflects EPA's updating of burden estimates for this collection based upon historical information on the number of responses anticipated for some ICs and the inclusion of an additional prior ICR titled, ''Exemptions of Certain Plant Incorporated Protectants (PIPs) Derived from Newer Technologies'' (OMB Control No. 2070–0214; Rulemaking RIN–2070–AK54). Based upon revised estimates and the inclusion of an additional IC, the total number of responses anticipated across categories has increased by 32,627 with a corresponding increase in the associated burden hours. While burden hours increased, the annual total costs to respondents have decreased due to a program change to reflect updated guidance on the calculation of overhead in fully loaded wages. This change is the result of a program change.

IV. What is the next step in the process for this ICR?

EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval pursuant to 5 CFR 1320.12. EPA will issue another **Federal Register** document pursuant to 5 CFR 1320.5(a)(1)(iv) to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB. If you have any questions about this ICR or the approval process, please contact the person listed under **FOR FURTHER INFORMATION CONTACT**.

Authority: 44 U.S.C. 3501 *et seq.*

Dated: November 27, 2025.

Nancy B. Beck,

Principal Deputy Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

[FR Doc. 2025–21758 Filed 12–1–25; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL MARITIME COMMISSION

[Docket No. 25–07]

PKDC, LLC, Complainant v. COSCO Shipping Lines Co., Ltd., Respondent; Notice of Filing of Amended Complaint

Notice is given that an amended complaint has been filed with the Federal Maritime Commission (the ''Commission'') by PKDC, LLC (the ''Complainant'') against COSCO Shipping Lines Co., Ltd. (the ''Respondent''). Complainant states that the Commission has subject-matter jurisdiction over this complaint pursuant to 46 U.S.C. 40101 *et seq.,* and personal jurisdiction over Respondent as an ocean common carrier, as defined in 46 U.S.C. 40102(7), (9), and (18).

Complainant is a limited liability company existing under the laws of the state of Colorado with its principal place of business in Denver, Colorado.

Complainant identifies Respondent as a corporation organized under the laws of the People's Republic of China with its corporate headquarters in Shanghai, China, whose agent in the United States is COSCO Shipping Lines (North America) Inc. with its principal place of business in Secaucus, New Jersey.

Complainant alleges that Respondent violated 46 U.S.C. 41102(c) and 46 CFR 545.5. Complainant alleges these violations arose from the assessment of unjust and unreasonable demurrage and detention charges due to circumstances outside the control of Complainant, 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/25-07/.* This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding judge shall be issued by May 6, 2026, and the final decision of the Commission shall be issued by November 20, 2026.

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

Served: November 28, 2025.

David Eng,

Secretary.

[FR Doc. 2025–21770 Filed 12–1–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–N–4731]

Increasing Access to Nonprescription Drugs; Request for Information

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for information.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing a request for information from interested parties and the public to share their perspectives with FDA on how to increase access to nonprescription drugs. The Agency intends to use the information submitted to inform plans for a public meeting intended to be held in calendar year 2026.

DATES: Either electronic or written comments on the notice must be submitted by February 2, 2026.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of February 2, 2026. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://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