Filing Party: Wayne Rohde, Cozen O'Connor.

Synopsis: The amendment would add a new Article 5.1(b)(ii) to provide for a structural provision of slots. It would also revise Article 9, including the addition of a new Article 9.5. It updates address and contact information in Article 3 and 11, and revises the governing law (Article 10) and arbitration (Article 13). The amendment also restates the agreement.

Proposed Effective Date: 8/1/2026.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/809.

Agreement No.: 201436–006.

Agreement Name: MSC/ZIM Cooperative Working Agreement.

Parties: Mediterranean Shipping Company S.A.; and Zim Integrated Shipping Services.

Filing Party: Wayne Rohde, Cozen O'Connor.

Synopsis: The amendment would revise the geographic scope of the Agreement by deleting Taiwan and adding India.

Proposed Effective Date: 7/30/2026.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/86581.

Agreement No.: 201471.

Agreement Name: Maersk/Hapag-Lloyd NAE Space Charter.

Parties: Maersk A/S; and Hapag Lloyd AG.

Filing Party: Wayne Rohde, Cozen O'Connor.

Synopsis: The agreement would authorize the chartering of space in the trade between ports on the U.S. East Coast, on the one hand, and ports in Colombia and Panama, on the other hand.

Proposed Effective Date: 8/1/2026.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/92676.

Dated: June 18, 2026.

Jennifer Everling,

Assistant Secretary.

[FR Doc. 2026–12630 Filed 6–23–26; 8:45 am]

BILLING CODE 6730–02–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (Act) (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the applications are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at *https://www.federalreserve.gov/foia/request.htm.* Interested persons may express their views in writing on the standards enumerated in paragraph 7 of the Act.

Comments received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Benjamin W. McDonough, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than July 9, 2026.

A. Federal Reserve Bank of St. Louis (Holly A. Rieser, Senior Manager) P.O. Box 442, St. Louis, Missouri 63166–2034. Comments can also be sent electronically to *Comments.applications@stls.frb.org:*

1. *John H. Garrett, individually and as trustee of each of the Filip J. Garrett Bank Trust u/a/d March 20, 2024, Sara E. Garrett Bank Trust u/a/d March 20, 2024, John H. Garrett Bank Trust u/a/d March 20, 2024, and Ida M. Garrett Bank Trust u/a/d March 20, 2024, all of Southport, North Carolina; Jeffrey S. Scott, individually and as trustee of each of the Hayden Scott Trust created under the Sharon K. Garrett Revocable Trust dated November 1, 2000, as amended, and the Avery Scott Trust created under the Sharon K. Garrett Revocable Trust dated November 1, 2000, as amended, Tyler Scott, Shelby McCaffrey, Richard W. Scott, all of Purdy, Missouri; Sara E. Garrett, Ida M. Garrett, and Filip J. Garrett, all of Springfield, Missouri; as a group acting in concert, to acquire voting shares of Purdy Bancshares, Inc., Monett, Missouri, and thereby indirectly acquire voting shares of First State Bank of the Ozarks, Purdy, Missouri.*

B. Federal Reserve Bank of San Francisco (Keith Dudley, Vice President) 101 Market Street, San Francisco, California 94105–1579. Comments can also be sent electronically to *SF.Supervision.Comments.Applications@sf.frb.org:*

1. *The Stanton Washington Trust Bank Voting Trust,* Peter F. Stanton, as trustee, both of Spokane, Washington; to acquire voting shares of W.T.B. Financial Corporation, and thereby indirectly acquire voting shares of Washington Trust Bank, both of Spokane, Washington.

Board of Governors of the Federal Reserve System.

Yao-Chin Chao,

Associate Secretary of the Board.

[FR Doc. 2026–12697 Filed 6–23–26; 8:45 am]

BILLING CODE P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–XXXX; Docket No. 2026–0232; Sequence No. 1]

Information Collection; Accessibility Conformance Report (ACR) Repository

AGENCY: Office of Governmentwide Policy, General Services Administration (GSA).

ACTION: Notice; request for comments.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve a new information collection requirement regarding the Accessibility Conformance Report (ACR) Repository.

DATES: Submit comments on or before August 24, 2026.

ADDRESSES:

Submit comments identified by Information Collection 3090–XXXX; Accessibility Conformance Report (ACR) Repository to *https://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching for ''Information Collection 3090–XXXX; Accessibility Conformance Report (ACR) Repository''. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 3090–XXXX; Accessibility Conformance Report (ACR) Repository''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–XXXX; Accessibility Conformance

Report (ACR) Repository'' on your attached document. If your comment cannot be submitted using *regulations.gov,* call or email the points of contact in the **FOR FURTHER INFORMATION CONTACT** section of this document for alternate instructions.

Instructions: Please submit comments only and cite Information Collection 3090–XXXX; Accessibility Conformance Report (ACR) Repository, in all correspondence related to this collection. Comments received generally will be posted without change to *regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *regulations.gov,* approximately two-to-three business days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Andrew Nielson, Director of the Government-wide IT Accessibility Program, Office of Government-wide Policy, GSA, at telephone 202–330–3036 or via email to *andrew.nielson@gsa.gov* for clarification of content.

SUPPLEMENTARY INFORMATION:

A. Purpose

GSA's Office of Governmentwide Policy (OGP) is in the process of developing an application to facilitate and provide a centralized repository of Accessibility Conformance Reports (ACRs) for information and communication technology (ICT) products.

Our ACR Repository is also in response to a requirement from OMB in M–24–08 to ''explore options for establishing a standardized accessibility conformance reporting process for government procurement of ICT, which should include a central repository of vendor accessibility conformance reports.''

Federal agencies often require submission of ACRs as part of quotes/bids in response to procurement solicitations. We intend to encourage product vendors to list/upload their ACRs into the repository to make it easier and more efficient for Federal employees involved in the acquisition and implementation of ICT and interested members of the public to locate ACRs when considering accessibility in the procurement process (as required by Section 508 and in the Federal Acquisition Regulation).

B. Annual Reporting Burden

There is no obligation for product owners to submit ACRs for any product. The total number of annual responses would be at the product owner's discretion.

Vendor Admin Respondents: 2500.
Vendor User Respondents: 1500.
Hours per Vendor Profile Creation: 10 mins.
Hours per Vendor User Creation: 3 mins.
ACRs per Vendor Account: 2.
Total ACRs Uploaded: 5000.
Hours per ACR Upload: 5 mins.
Total Burden Hours: 907 hours.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary, whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies Of Proposals: Requesters may obtain a watermarked ''DRAFT'' copy of the supporting statement via the Federal eRulemaking portal at *regulations.gov* by searching for Docket ID ''GSA–GSA–2026–0232.'' Select the document titled ''Supporting Statement: 3090–XXXX, Accessibility Conformance Report (ACR) Repository—DRAFT'' located under Supporting and Related Material.''

Richard Speidel,

Deputy Chief Data Officer, General Services Administration.

[FR Doc. 2026–12667 Filed 6–23–26; 8:45 am]

BILLING CODE 6820–WY–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–D–1494]

International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Effectiveness of Anthelmintics: General Recommendations; 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 final guidance for industry (GFI) #90 entitled ''Effectiveness of Anthelmintics: General Recommendations.'' This guidance has been developed for veterinary use by the International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH). The objective of this guidance is to provide study design recommendations that will facilitate the universal acceptance of the generated effectiveness data to fulfill the national/regional requirements for anthelmintic drugs in animal species.

DATES: The announcement of the guidance is published in the **Federal Register** on June 24, 2026.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

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 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 *https://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):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, 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.''