Obligation to Respond: Required to obtain or retain benefits. The statutory authority for the information collection requirements is contained in sections 4(i), 4(j), 13, 201(b), 254, 257, 301, 303, 316, and 332 of the Communications Act of 1934, as amended, 47 U.S.C. 154(i), 154(j), 163, 201(b), 254, 257, 301, 303, 316, 332, section 60504 of the Infrastructure Investment and Jobs Act, Public Law 117–58, 135 Stat. 429 (2021), and section 904 of the Consolidated Appropriations Act, 2021, Public Law 116–260, 134 Stat. 1182 (2020), as amended.

Total Annual Burden: 983,493 hours.

Total Annual Cost: $4,250,000.

Needs and Uses: This information collection pertains to the Empowering Broadband Consumers Through Transparency, Report and Order and Further Notice of Proposed Rulemaking, published at 87 FR 76959 (Dec. 16, 2022) (Broadband Label Order). The information will be used to implement section 60504(a) of the Infrastructure Investment and Jobs Act (Infrastructure Act). The Infrastructure Act, in relevant part, directed the Commission ''[n]ot later than 1 year after the date of enactment of th[e] Act, to promulgate regulations to require the display of broadband consumer labels, as described in the Public Notice of the Commission issued on April 4, 2016 (DA 16–357), to disclose to consumers information regarding broadband internet access service plans.'' Further, the Infrastructure Act required that the label ''include information regarding whether the offered price is an introductory rate and, if so, the price the consumer will be required to pay following the introductory period.''

On January 27, 2022, the Commission released a Notice of Proposed Rulemaking, published at 87 FR 6827 (Feb. 7, 2022), initiating a proceeding to implement section 60504 of the Infrastructure Act. Specifically, the Commission proposed to require that broadband internet access service providers (ISPs or providers) display, at the point of sale, labels that disclose to consumers certain information about prices, introductory rates, data allowances, broadband speeds, and management practices, among other things.

On November 14, 2022, the Commission adopted the Broadband Label Order requiring ISPs to display a new broadband label to help consumers comparison shop among broadband services, thereby implementing section 60504 of the Infrastructure Act. Specifically, the Commission required ISPs to display, at the point of sale, a broadband consumer label containing critical information about the provider's service offerings, including information about pricing, introductory rates, data allowances, performance metrics, and whether the provider participates in the Affordable Connectivity Program (ACP). The Commission required that ISPs display the label for each stand-alone broadband internet access service they currently offer for purchase, and that the label link to other important information such as network management practices, privacy policies, and other educational materials. Consistent with the Infrastructure Act, the label adopted for fixed and mobile broadband internet access service is similar to the two voluntary labels the Commission approved in 2016, with certain modifications. The label resembles the well-known nutrition labels that consumers have come to rely on when shopping for food products.

In addition to label content, the Commission adopted requirements for the label's format and display location to ensure consumers can make side-by-side comparisons of various service offerings from an individual provider or from alternative providers—something essential for making informed decisions. Labels must be displayed on providers' websites and at alternate sales channels such as retail locations and over the phone. The label must be accessible for people with disabilities and for non-English speakers. Labels must also be available via a customer's online account portal. ISPs shall maintain an archive of all labels for a period of no less than two years from the time the service plan reflected in the label is no longer available for purchase by a new subscriber and the provider has removed the label from its website or alternate sales channels. In addition, third parties will be able to easily analyze information contained in the labels and help consumers with their purchase decisions, as providers are required to make the label content available in a machine-readable format on their websites. Finally, the Commission adopted a label template that all ISPs are required to display at the point of sale. This label establishes the formatting and content of all requirements adopted in the Broadband Label Order.

Federal Communications Commission.

Aleta Bowers,

Federal Register Liaison Officer, Office of the Secretary.

[FR Doc. 2026–13220 Filed 6–30–26; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

Notice of Agreement Filed

The Commission hereby gives notice of filing of the following agreement under the Shipping Act of 1984. Interested parties may submit comments, relevant information, or documents regarding the agreement to the Secretary by email at *Secretary@fmc.gov,* or by mail, Federal Maritime Commission, 800 North Capitol Street Washington, DC 20573. Comments will be most helpful to the Commission if received within 12 days of the date this notice appears in the **Federal Register**, and the Commission requests that comments be submitted within 7 days on agreements that request expedited review. Copies of the agreement are available through the Commission's website (*www.fmc.gov*) or by contacting the Office of General Counsel at (202)–523–5740 or *GeneralCounsel@fmc.gov.*

Agreement No.: 201472.

Agreement Name: Mitsui O.S.K. Lines Ltd./Grimaldi Deep Sea S.p.A./Grimaldi Euromed S.p.A. South America Space Charter Agreement.

Parties: Mitsui O.S.K. Lines Ltd.; and Grimaldi Deep Sea S.P.A. and Grimaldi Euromed S.p.A. (acting as a single party).

Filing Party: Rebecca Fenneman, Jeffrey Fenneman Law and Strategy, PLLC.

Synopsis: The Agreement authorizes the parties to charter space for the carriage of cargo including, but not limited to any and all classes of roll-on/roll-off cargo to/from one another in the trade on an ''as needed/as available'' basis.

Proposed Effective Date: 6/19/2026.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/Document/131579.

Dated: June 29, 2026.

Jennifer Everling,

Assistant Secretary.

[FR Doc. 2026–13291 Filed 6–30–26; 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 16, 2026.

A. Federal Reserve Bank of Dallas (Lindsey Wieck, Director, Mergers & Acquisitions) 2200 North Pearl Street, Dallas, Texas 75201–2272. Comments can also be sent electronically to *Comments.applications@dal.frb.org:*

1. *Austin Family Trust B, Jeff Austin, Jr. as trustee, both of Jacksonville, Texas; JML Trust, Mary Margaret Austin, as trustee, both of Jacksonville, Texas; and LAPA Trust, Carole Leigh Austin Mattson, as trustee, both of Georgetown, Texas;* to join the Austin/Chapman Family Control Group, a group acting in concert, to acquire voting shares of Austin Bancorp, Inc., and thereby indirectly acquire voting shares of Austin Bank, Texas National Association, both of Jacksonville, Texas.

2. *Austin Family Trust B, Jeff Austin, Jr. as trustee, both of Jacksonville, Texas;* to acquire voting shares of Capital Bancorp, Inc., Jacinto City, Texas, and thereby indirectly acquire voting shares of Capital Bank, Houston, Texas.

In addition,*JML Trust, Mary Margaret Austin, as trustee, both of Longmont, Colorado; LAPA Trust, Carole Leigh Austin Mattson, as trustee, both of Littleton, Colorado; Jessica Leigh Neill Swinnea, Chandler, Texas; and Austin Kyle Neill, Dallas, Texas;* to join the Austin/Chapman Family Control Group, a group acting in concert, to retain voting shares of Capital Bancorp, Inc., and thereby indirectly retain voting shares of Capital Bank.

3. *Austin Family Trust B, Jeff Austin, Jr. as trustee, both of Jacksonville, Texas;* to join the Austin/Chapman Family Control Group, a group acting in concert, to acquire voting shares of Athens, TX Bancshares, Inc., and thereby indirectly acquire voting shares of First State Bank, both of Athens, Texas.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026–13287 Filed 6–30–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–N–6869]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medication Guides for Prescription Drug Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by July 31, 2026.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function. The OMB control number for this information collection is 0910–0393. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Medication Guide Requirements for Prescription Drug Product Labeling

OMB Control Number 0910–0393—Reinstatement with change

This information collection supports FDA regulations pertaining to the distribution of patient labeling, called Medication Guides, for human prescription drug and biological products used primarily on an outpatient basis, and required for products that pose a serious and significant public health concern. Agency regulations codified in part 208 (21 CFR part 208): *Medication Guides for Prescription Drug Products* set forth general requirements applicable to both content and format elements for Medication Guides, as well as provide for exemption and deferral requests. Medication Guides provide patients with important information about drug products, including the drug's approved uses, contraindications, adverse drug reactions, cautions for specific populations, and are required in accordance with applicable regulations.

To assist consumers and industry with understanding regulatory requirements applicable to, and the purpose of, Medication Guides, we have developed resources and made them available on our website at *https://www.fda.gov/drugs/fdas-labeling-resources-human-prescription-drugs/patient-labeling-resources#medication-guides.* Among the resources, we include the guidance document titled *Medication Guides—Distribution Requirements and Inclusion in Risk Evaluation and Mitigation Strategies (REMS)* (November 2011), (available at *https://www.fda.gov/media/79776/download*), as well as a discussion of the distinction between Medication Guides and Consumer Medication information. The regulations in 21 CFR part 208, the associated guidance document, and the informational resources are intended to enable patients to use medications most safely and effectively.

Included among the requirements in 21 CFR part 314 that govern applications for FDA approval to market a new drug is the submission of any Medication Guide (see 21 CFR part