assessment rate in the agreement, and also accordingly revises various figures set forth in Appendix 1.

Proposed Effective Date: 6/5/2026.

Location: https://www2.fmc.gov/FMC.Agreements.Web/Public/AgreementHistory/10164.

Agreement No.: 201436–005.

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 add the Dominican Republic to the geographic scope of the Agreement. The parties have requested expedited review.

Proposed Effective Date: 7/23/2026.

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

Dated: June 12, 2026.

Jennifer Everling,

Assistant Secretary.

[FR Doc. 2026–12136 Filed 6–16–26; 8:45 am]

BILLING CODE 6730–02–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

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 the BHC Act (12 U.S.C. 1842(c)).

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 17, 2026.

A. Federal Reserve Bank of Richmond (Brent B. Hassell, Assistant Vice President) P.O. Box 27622, Richmond, Virginia 23261. Comments can also be sent electronically to *Comments.applications@rich.frb.org:*

1. *Client First Holdings, LLC, Coeburn, Virginia;* to become a bank holding company by acquiring Miners Exchange Bank through a merger with a newly formed subsidiary, Client First Interim Bank, both of Coeburn, Virginia.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026–12209 Filed 6–16–26; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Notice of Board Meeting

DATES: June 23, 2026 at 11:00 a.m. ET.

ADDRESSES: Telephonic. Dial-in (listen only) information: Number: 1–202–599–1426, Code: 243 438 047#; or via web: *https://www.frtib.gov/.*

FOR FURTHER INFORMATION CONTACT: James Kaplan, Director, Office of External Affairs, (202) 864–7150.

SUPPLEMENTARY INFORMATION:

Board Meeting Agenda

Open Session

1. Approval of the May 28, 2026, Board Meeting Minutes
2. Monthly Reports
   (a) Participant Report
   (b) Investment Report
   (c) Legislative Report
3. Quarterly Reports
   (d) Vendor Risk Management

Authority: 5 U.S.C. 552b (e)(1).

Dated: June 12, 2026.

Dharmesh Vashee,

General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 2026–12133 Filed 6–16–26; 8:45 am]

BILLING CODE 6760–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Information collection notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the extension with change of the currently approved information collection ''AHRQ Research Reporting System (ARRS)'' OMB No. 0935–0122, reflecting a reduced annual reporting burden and expanded applicability to AHRQ-supported extramural research activities conducted through grants, contracts, and challenge competitions.

DATES: Comments on this notice must be received by August 17, 2026.

ADDRESSES: Written comments should be submitted to: Margie Shofer, Reports Clearance Officer, AHRQ, by email at *REPORTSCLEARANCEOFFICER@ahrq.hhs.gov.*

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Margie Shofer, AHRQ Reports Clearance Officer, (301) 427–1696, or by email at *REPORTSCLEARANCEOFFICER@ahrq.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

In 2007, AHRQ developed a systematic method for its grantees to report project progress, and important preliminary findings called the Grants Reporting System (GRS). In 2008, GRS was renamed AHRQ Research Reporting System (ARRS) and in addition to reporting on grant progress, also supported reporting on jobs created under contracts pursuant to the American Recovery and Reinvestment Act of 2009. ARRS is still being used to

track grant progress and may also support administrative progress reporting for AHRQ-funded research activities conducted under contracts and challenge competition awards, as applicable. Such information may be entered by award recipients, contractors, or authorized AHRQ officials, as appropriate. ARRS is no longer used for contract job creation.

The system addressed the shortfalls in the previous reporting process and established a consistent and comprehensive reporting solution for AHRQ. The ARRS provides a centralized repository of AHRQ-supported research progress and administrative project information across grants, and where applicable, research contracts and challenge competition awards, that can be used to support initiatives within the Agency. This includes future research planning and support to administration activities such as performance monitoring, budgeting, knowledge transfer as well as strategic planning.

This Project seeks to answer the following research questions:

(1) What progress has been demonstrated across AHRQ-funded research activities and related funding mechanisms, including grants, research contracts, and challenge completion awards, and how can the resulting progress and output data be systematically leveraged to inform future research planning and to support core administrative functions, including performance monitoring, budgeting, knowledge transfer, and strategic planning?

This Project has the following goals:

(1) To promote the transfer of critical information more frequently and efficiently and enhance the Agency's ability to support research designed to improve the outcomes and quality of health care, reduce its costs, and broaden access to effective services

(2) To increase the efficiency of the Agency in responding to ad-hoc information requests

(3) To support Executive Branch requirements for increased transparency and public reporting

(4) To establish a consistent approach throughout the Agency for information collection regarding research progress and a systematic basis for oversight and for facilitating potential collaborations among grantees

(5) To decrease the inconvenience and burden on grantees of unanticipated ad-hoc requests for information by the Agency in response to particular (one-time) internal and external requests for information

The information is being collected by AHRQ, pursuant to its statutory authority to conduct and support research on healthcare and on systems for the delivery of such care, including activities with respect to the quality, effectiveness, efficiency, appropriateness and value of healthcare services and with respect to quality measurement and improvement, and database development. 42 U.S.C. 299a(a)(1) and (8). Minor revisions to the scope of respondents are being proposed to clarify applicability to AHRQ-funded research contracts and challenge competition recipients; no substantive revisions to the core reporting content are being proposed.

Method of Collection

To achieve the goals of this project the following data collections will be implemented:

AHRQ Research Reporting System (ARRS)—Award recipients, contractors, and authorized AHRQ official, as applicable, use the ARRS system to report project progress and important preliminary findings for AHRQ-funded research activities. Reporting frequency varies based on award type and programmatic need. All users access the ARRS system through a secure online interface which requires user authentication. When status reports are due AHRQ notifies designated reporting officials via email.

Estimated Annual Respondent Burden

Exhibit 1 shows the estimated annualized burden hours for the respondents. The estimated number of respondents is 450 a year, a decrease from the last Information Collection Request, which estimated 500 reports to be collected in a year. This revised amount is based on the current number of 210 active reports and adjusted upwards to account for any new grants, research contracts, or challenge competition awards AHRQ may award in the future.

Grantees will take an estimated 30 minutes to enter the necessary data into the ARRS. Frequency of reporting varies from monthly to twice a year. Based on that, the total annualized burden hours are estimated to be 225 hours.

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS

Form Name	Number of respondents	Number of responses per respondent	Hours per response	Total burden hours
ARRS Data Entry	450	1	30/60	225
Total	450	N/A	N/A	225

Exhibit 2 shows the estimated annualized cost burden for the respondents. The total estimated cost burden for respondents is $26,725.50.

EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN

Form Name	Total burden hours	Average hourly wage rate *	Adjusted hourly wage rate **	Total cost burden
ARRS Data Entry	225	$59.39	$118.78	$26,725.50
Total	225	N/A	N/A	26,725.50

"National Compensation Survey: Occupational Wages in the United States, May 2025," U.S. Department of Labor, Bureau of Labor Statistics, *http://www.bls.gov/oes/current/oes_nat.htm#29-0000.*

* Based upon the hourly 75th percentile wage for Healthcare Practitioner and Technical Occupations (29–0000) since most responders are healthcare clinicians.

** The Adjusted Hourly Rate was estimated at 200% of the hourly wage.

Request for Comments

In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3520, comments on AHRQ's information collection are requested with regard to any of the following: (a) whether the proposed collection of information is necessary for the proper performance of AHRQ's health care research and health care information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Dated: June 12, 2026

Jeffrey Toven,

Executive Officer.

[FR Doc. 2026–12174 Filed 6–16–26; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–1123]

Andrew Jonathan Morgan: Final Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is issuing an order under the Federal Food, Drug, and Cosmetic Act (FD&C Act) debarring Andrew Jonathan Morgan for a period of 5 years from importing or offering for import any drug into the United States. FDA bases this order on a finding that Mr. Morgan was convicted of a felony under federal law. The factual basis supporting Mr. Morgan's conviction, as described below, is conduct relating to the importation into the United States of a drug or controlled substance. Mr. Morgan was given notice of the proposed debarment and was given an opportunity to request a hearing to show why he should not be debarred. As of April 29, 2026 (30 days after receipt of the notice), Mr. Morgan had not responded. Mr. Morgan's failure to respond and request a hearing constitutes a waiver of his right to a hearing concerning this matter.

DATES: This order is applicable June 17, 2026.

ADDRESSES: Any application by Mr. Morgan for termination of debarment under section 306(d)(1) of the FD&C Act (21 U.S.C. 335a(d)(1)) may be submitted at any time as follows:

Electronic Submissions

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. An application submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your application will be made public, you are solely responsible for ensuring that your application 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 application, that information will be posted on *https://www.regulations.gov.*
- If you want to submit an application with confidential information that you do not wish to be made available to the public, submit the application as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

- *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 a written/paper application submitted to the Dockets Management Staff, FDA will post your application, as well as any attachments, except for information submitted, marked, and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All applications must include the Docket No. FDA–2026–N–1123. Received applications will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit an application with confidential information that you do not wish to be made publicly available, submit your application only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of your application. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852 between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500. Publicly available submissions may be seen in the docket.

FOR FURTHER INFORMATION CONTACT: Jaime Espinosa, Division of Field Enforcement, Office of Field Regulatory Operations, Office of Inspections and Investigations, Food and Drug Administration, 240–402–8743, or *debarments@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

Section 306(b)(1)(D) of the FD&C Act permits debarment of an individual from importing or offering for import any drug into the United States if FDA finds, as required by section 306(b)(3)(C) of the FD&C Act, that the individual has been convicted of a felony for conduct relating to the importation into the United States of any drug or controlled substance.