public is encouraged to contact OPP at (202) 502–6595 or *OPP@ferc.gov.*

Dated: May 28, 2025.

Debbie-Anne A. Reese,

Secretary.

[FR Doc. 2025–10099 Filed 6–3–25; 8:45 am]

BILLING CODE 6717–01–P

EXPORT-IMPORT BANK

[Public Notice: EIB–2025–0005]

Application for Final Commitment for a Long-Term Loan or Financial Guarantee in Excess of $100 Million: AP089541XX

AGENCY: Export-Import Bank of the United States.

ACTION: Notice.

SUMMARY: This Notice is to inform the public, in accordance with the Export-Import Bank Act of 1945, as amended, the Export-Import Bank of the United States (''EXIM'') has received an application for final commitment for a long-term loan or financial guarantee in excess of $100 million. Comments received within the comment period specified below will be presented to the EXIM Board of Directors prior to final action on this Transaction.

DATES: Comments must be received on or before June 30, 2025 to be assured of consideration before final consideration of the transaction by the Board of Directors of EXIM.

ADDRESSES: Comments may be submitted through *Regulations.gov* at *www.regulations.gov.* To submit a comment, enter *EIB–2025–0005* under the heading ''Enter Keyword or ID'' and select Search. Follow the instructions provided at the Submit a Comment screen. Please include your name, company name (if any) and *EIB–2025–0005* on any attached document.

SUPPLEMENTARY INFORMATION:

Reference: AP089541XX.

Purpose and Use:

Brief description of the purpose of the transaction: To support the export of U.S.-manufactured commercial aircraft and spare engines to Angola.

Brief non-proprietary description of the anticipated use of the items being exported: To provide passenger and cargo air transport between Angola and other countries.

To the extent that EXIM is reasonably aware, the item(s) being exported may be used to produce exports or provide services in competition with the exportation of goods or provision of services by a United States industry.

Parties:

Principal Supplier: The Boeing Company and GE Aerospace.

Obligor: TAAG Angola Airlines.

Guarantor(s): The African Export-Import Bank.

Description of Items Being Exported: Boeing commercial jet aircraft and GE spare engines.

Information on Decision: Information on the final decision for this transaction will be available in the ''Board Agenda and Meeting Minutes'' on *https://www.exim.gov/news/meeting-minutes.*

Confidential Information: Please note that this notice does not include confidential or proprietary business information; information which, if disclosed, would violate the Trade Secrets Act; or information which would jeopardize jobs in the United States by supplying information that competitors could use to compete with companies in the United States.

Authority: Section 3(c)(10) of the Export-Import Bank Act of 1945, as amended (12 U.S.C. 635a(c)(10)).

Deidre Hodge,

Assistant Corporate Secretary.

[FR Doc. 2025–10154 Filed 6–3–25; 8:45 am]

BILLING CODE 6690–01–P

FEDERAL MARITIME COMMISSION

Notice of Agreements Filed

The Commission hereby gives notice of the filing of the following agreements under the Shipping Act of 1984. Interested parties may submit comments, relevant information, or documents regarding the agreements 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 agreements are available through the Commission's website (*www.fmc.gov*) or by contacting the Office of Agreements at (202)–523–5793 or *tradeanalysis@fmc.gov.*

Agreement No.: 201436–003.

Agreement Name: MSC/ZIM Cooperative Working Agreement.

Parties: Mediterranean Shipping Company SA; and ZIM Integrated Shipping Services Ltd.

Filing Party: Wayne Rohde, Cozen O'Connor.

Synopsis: The Amendment deletes Indonesia from the geographic scope of the agreement and makes adjustments to Article 14.3 in light of increased demand resulting from changes in trade conditions.

Proposed Effective Date: 5/26/2025.

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

Agreement No.: 201436–004.

Agreement Name: MSC/ZIM Cooperative Working Agreement.

Parties: Mediterranean Shipping Company SA; and ZIM Integrated Shipping Services Ltd.

Filing Party: Wayne Rohde, Cozen O'Connor.

Synopsis: The Amendment would add the Bahamas, Mexico, Jamaica, Sri Lanka and Indonesia to the geographic scope of the Agreement.

Proposed Effective Date: 7/12/2025.

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

Dated: May 30, 2025.

Alanna Beck,

Federal Register Alternate Liaison Officer.

[FR Doc. 2025–10160 Filed 6–3–25; 8:45 am]

BILLING CODE 6730–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10277, CMS–265–11 and CMS–10916]

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 August 4, 2025.

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**).

CMS–10277—Medicare and Medicaid Programs: Conditions of Participation for Hospices

CMS–265–11—Independent Renal Dialysis Facility Cost Report

CMS–10916—13th SOW QIN–QIO and AI/AN Advancing Healthcare Quality through Technology (AHQT) Readiness Assessment

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 Collections

1. *Type of Information Collection Request:* Reinstatement with change of a previously approved collection; *Title of Information Collection:* Medicare and Medicaid Programs: Conditions of Participation for Hospices; *Use:* Under the Medicare program, eligible beneficiaries may receive covered services in a hospice, provided that certain requirements are met by the hospice. Hospice care means a comprehensive set of services identified and coordinated by an interdisciplinary group to provide for the physical, psychosocial, spiritual and emotional needs of a terminally ill patient and/or family members, as delineated in a specific patient plan of care.

The information collection requirements (ICRs) described herein are needed to implement the Medicare Conditions of Participation (CoPs) for Medicare-participating hospices. The CoPs help assure an adequate level of patient health and safety in participating hospices and help ensure that Medicare hospice eligibility requirements are being met. CMS originally published the Hospice Conditions of Participation on June 5, 2008 (hereinafter ''*2008 Final Rule*''). The regulations containing the information collection requirements are located at 42 CFR part 418 of the Code of Federal Regulations, Subparts B, C and D.

This is a reinstatement of the information collection request that expired on March 31, 2024. The previous iteration of this OMB Control Number: 0938–1067 (approved March 23, 2021) had an annual burden of 3,639,215 hours and annual costs of $273,001,454. For this requested reinstatement, with changes, the total annual burden hours for industry is 4,032,329 hours and the annual burden costs are $350,449,922. The 10.8% increase in hours is primarily due to the increase in the number of hospices since the last iteration.

Since the last reinstatement was approved in March 2021, CMS revised one of the hospice CoPs at 42 CFR 418.76 in the proposed rule, *Medicare Program: FY 2022 Hospice Wage Index and Payment Rate Update, Hospice Conditions of Participation Updates, Hospice and Home Health Quality Reporting Program Requirements* published on April 14, 2021 (86 FR 19700). As CMS addressed in the final rule (CMS–1754–F) published on August 4, 2021 (86 FR 42528), the comments received supported the proposed revisions and did not require any changes to the original burden estimates in this PRA package. This reinstatement incorporates the policy changes made to Section 418.76 through this rule and updates the associated burden estimates based on the original assumptions.

In November 2021, CMS required hospices to develop policies and procedures as a CoP to ensure all staff were fully vaccinated and the burden requirements were detailed in OMB Control Number: 0938–0266. However, CMS removed this requirement and related burden for hospices (and other facilities) in June 2023. *Form Number:* CMS–10277 (OMB control number: 0938–1067); *Frequency:* Yearly; *Affected Public:* Private sector (Business or other for-profit and Not-for-profit institutions); *Number of Respondents:* 7,356; *Total Annual Responses:* 9,209,893; *Total Annual Hours:* 4,032,329. (For policy questions regarding this collection contact Claudia Molinar at 410–786–8445.)

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Independent Renal Dialysis Facility Cost Report; *Use:* Under the authority of sections 1815(a) and 1833(e) of the Act, CMS requires that providers of services participating in the Medicare program submit information to determine costs for health care services rendered to Medicare beneficiaries. CMS requires that providers follow reasonable cost principles under 1861(v)(1)(A) of the Act when completing the Medicare cost report (MCR). Regulations at 42 CFR 413.20 and 413.24 require that providers submit acceptable cost reports on an annual basis and maintain sufficient financial records and statistical data, capable of verification by qualified auditors.

ESRD facilities participating in the Medicare program submit these cost reports annually to report cost and statistical data used by CMS to

determine reasonable costs incurred for furnishing dialysis services to Medicare beneficiaries and to effect the year-end cost settlement for Medicare bad debts. *Form Number:* CMS–265–11 (OMB control number: 0938–0236); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for-profits, State, Local, or Tribal Governments); *Number of Respondents:* 7,329; *Total Annual Responses:* 7,329; *Total Annual Hours:* 483,714. (For questions regarding this collection contact Keplinger, Jill C. at 410–786–4550.)

3. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* 13th SOW QIN–QIO and AI/AN Advancing Healthcare Quality through Technology (AHQT) Readiness Assessment; *Use:* This is a new information collection request. The Quality Improvement Network—Quality Improvement Organization (QIN–QIO) program and American Indian/Alaska Native (AI/AN) program assists providers/practices with high-quality, hands-on quality improvement assistance toward meeting their needs, and the healthcare quality and safety goals for beneficiaries. The purpose of this new information collection within these programs is to assess the readiness of participating nursing homes, hospitals, outpatient clinical practices, and AIAN facilities to access, share, and use data electronically for quality improvement and quality reporting. Use of health information technology (HIT) is imperative to assess, monitor, and improve healthcare quality, patient safety, and care coordination.

Many providers/practices continue to lack basic knowledge and capacity to implement HIT to support data exchange between providers/practices, payers, and patients, and to use data for improving quality and outcomes. This ''digital divide'' creates burden for patients, families, caregivers, providers/practices and increases costs and administrative waste. This burden is disproportionate for underserved populations. Advancing the use of technology and using interoperable standards can reduce the overall cost and burden associated with data collection and supports communication across the care continuum and is an agency priority.

CMS has developed a 41-item Assessment of Health care Quality Technical Readiness (AHQT) for use with participating providers/practices under the QIN–QIO 13th SOW. Provider/practice burden associated with the collection and reporting of quality measurement data has historically been a pain point for the QIN–QIO and AI/AN programs, especially in outpatient clinical practices and critical access hospitals; this burden has been a barrier to both achievement of quality improvement contract goals and proper evaluation of their impact. The results of the assessment will be used to determine which providers/practices may benefit from participation in a technical assistance pilot specific during the QIN–QIO 13th SOW intended to advance provider/practice capacity for engaging in quality improvement and reporting activities facilitated by HIT. *Form Number:* CMS–10916 (OMB control number: 0938–NEW); *Frequency:* Occasionally; *Affected Public:* Private Sector (Business or other for-profit and Not-for-profit institutions); *Number of Respondents:* 53,000; *Total Annual Responses:* 10,600; *Total Annual Hours:* 10,600. (For policy questions regarding this collection contact Geoffrey Berryman at 410–299–7390).

William N. Parham, III,

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

[FR Doc. 2025–10112 Filed 6–3–25; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Cancellation of Meeting

Notice is hereby given of the cancellation of the Center for Scientific Review Special Emphasis Panel Program Project National Center for Biomedical Imaging and Bioengineering, June 02, 2025, 10:00 a.m. to June 04, 2025, 5:00 p.m., National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD, 20892 which was published in the **Federal Register** on May 9, 2025, 90 FR 19725, Doc. No. 2025–08184.

This notice is being amended to announce that the meeting is cancelled.

Dated: May 30, 2025.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2025–10143 Filed 6–3–25; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Neuromodulation and Imaging of Neuronal Circuits.

Date: June 30–July 1, 2025.

Time: 9:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Pablo Miguel Blazquez Gamez, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 435–1042, *pablo.blazquezgamez@nih.gov.*

Name of Committee: Biological Chemistry and Macromolecular Biophysics Integrated Review Group; Chemical Synthesis and Biosynthesis Study Section.

Date: June 30–July 1, 2025.

Time: 9:30 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Shan Wang, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 496–4390, *shan.wang@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: May 30, 2025.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2025–10144 Filed 6–3–25; 8:45 am]

BILLING CODE 4140–01–P