information is not publicly available, *e.g.,* CBI or other information whose disclosure is restricted by statute. Certain other materials, such as copyrighted material, are publicly available only in hard copy. Publicly available docket materials are available either electronically in *www.regulations.gov* or in hard copy at the ORD Docket in EPA's Headquarters Docket Center.

Wayne Cascio,

Director, Center for Public Health and Environmental Assessment, Office of Research and Development.

[FR Doc. 2022–27939 Filed 12–30–22; 8:45 am]

BILLING CODE 6560–50–P

EXPORT-IMPORT BANK

Updated Intent To Conduct a Detailed Economic Impact Analysis

AGENCY: Export-Import Bank.

ACTION: Notice.

SUMMARY: This notice is to inform the public that the Export-Import Bank of the United States has received has received a request to increase the financed amount for a previously notified application (FR Doc. 2022–19164). The application is now for a $99.7 million direct loan to support the export of approximately $63.88 million in U.S. equipment and services to upgrade and expand an oil refinery in Indonesia. There has been no change in the expected output of the facility, and the supported U.S. exports will still enable the facility to increase production capacity of gasoline by 101,000 barrels per day and propylene by 225,000 tons per year. Available information indicates that the refined products will be consumed domestically, with negligible sales to other markets.

DATES: Comments are due 14 days from publication in the **Federal Register**.

ADDRESSES: Interested parties may submit comments on this transaction electronically at *www.regulations.gov,* or by email to *economic.impact@exim.gov*.

Scott Condren,

Sr. Policy Analyst.

[FR Doc. 2022–28495 Filed 12–30–22; 8:45 am]

BILLING CODE 6690–01–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0548; FR ID 121020]

Information Collection Being Reviewed by the Federal Communications Commission

AGENCY: Federal Communications Commission.

ACTION: Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act of 1995 (PRA), the Federal Communications Commission (FCC or Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid Office of Management and Budget (OMB) control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

DATES: Written comments shall be submitted on or before March 6, 2023. If you anticipate that you will be submitting comments but find it difficult to do so within the period of time allowed by this notice, you should advise the contacts below as soon as possible.

ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email: *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418–2918.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0548.

Title: Sections 76.1709 and 76.1620, Availability of Signals; Section 76.1614, Identification of Must-Carry Signals.

Type of Review: Extension without change of a currently approved collection.

Respondents: Businesses or other for-profit.

Number of Respondents and Responses: 4,103 respondents; 49,236 responses.

Estimated Time per Response: 0.5–1.0 hour.

Frequency of Response: Recordkeeping requirement, Third party disclosure requirement, On occasion reporting requirement.

Obligation to Respond: Voluntary.

Total Annual Burden: 24,618 hours.

Total Annual Cost: No cost.

Needs and Uses: 47 CFR 76.1709(a) states that the operator of every cable television system shall maintain for public inspection a file containing a list of all broadcast television stations carried by its system in fulfillment of the must-carry requirements. Such list shall include the call sign; community of license, broadcast channel number, cable channel number, and in the case of a noncommercial educational broadcast station, whether that station was carried by the cable system on March 29, 1990.

47 CFR 76.1614 and 47 CFR 76.1709(c) each state that a cable operator shall respond in writing within 30 days to any written request by any person for the identification of the signals carried on its system in fulfillment of the must-carry requirements. In addition, 47 CFR 76.1614 states that the required written response may be delivered by email, if the consumer used email to make the request or complaint directly to the cable operator, or if the consumer specifies email as the preferred delivery method in the request or complaint.

47 CFR 76.1620, pursuant to 47 U.S.C. 614(b)(7), states that if a cable operator authorizes subscribers to install additional receiver connections, but does not provide the subscriber with such connections, or with the equipment and materials for such connections, the operator shall notify such subscribers of all broadcast stations carried on the cable system which cannot be viewed via cable without a converter box and shall offer to sell or lease such a converter box to such subscribers. Such notification must be provided by June 2, 1993, and annually thereafter and to each new subscriber upon initial installation. The notice, which may be included in routine billing statements, shall identify the signals that are unavailable without

an additional connection, the manner for obtaining such additional connection and instructions for installation. 47 CFR 76.1600(a) provides that written information provided by cable operators to subscribers or customers pursuant to § 76.1620 may be delivered electronically by email to any subscriber who has not opted out of electronic delivery if the entity: (1) Sends the notice to the subscriber's or customer's verified email address; (2) Provides either the entirety of the written information or a weblink to the written information in the notice; and (3) Includes, in the body of the notice, a telephone number that is clearly and prominently presented to subscribers so that it is readily identifiable as an opt-out mechanism that will allow subscribers to continue to receive paper copies of the written material.

Note: These recordkeeping and notification requirements ensure that subscribers are aware of the broadcast stations carried in compliance with the Commission's cable must-carry rules, see 47 CFR 76.56.

Federal Communications Commission.

Kimberly Stewart,

Federal Register Liaison Officer, Office of the Secretary.

[FR Doc. 2022–28491 Filed 12–30–22; 8:45 am]

BILLING CODE 6712–01–P

GOVERNMENT ACCOUNTABILITY OFFICE

Request for Medicaid and CHIP Payment and Access Commission (MACPAC) Nominations

AGENCY: Government Accountability Office.

ACTION: Request for letters of nomination and resumes.

SUMMARY: The Children's Health Insurance Program Reauthorization Act of 2009 (CHIPRA) established MACPAC to review Medicaid and CHIP access and payment policies and to advise Congress on issues affecting Medicaid and CHIP. CHIPRA gave the Comptroller General of the United States responsibility for appointing MACPAC's members. The U.S. Government Accountability Office (GAO) is now accepting nominations for MACPAC appointments that will be effective May 2023. Nominations should be sent to the email address listed below. Acknowledgement of receipt will be provided within a week of submission.

DATES: Letters of nomination and resumes should be submitted no later than January 26, 2023, to ensure adequate opportunity for review and consideration of nominees prior to appointment.

ADDRESSES: Submit letters of nomination and resumes to *MACPACappointments@gao.gov.*

FOR FURTHER INFORMATION CONTACT: Susan Anthony at (312) 220–7666 or *anthonys@gao.gov* if you do not receive an acknowledgment or need additional information. For general information, contact GAO's Office of Public Affairs, (202) 512–4800.

Authority: 42 U.S.C. 1396.

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2022–27887 Filed 12–30–22; 8:45 am]

BILLING CODE 1610–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0973]

Revocation of Three Authorizations of Emergency Use of In Vitro Diagnostic Devices for Detection and/or Diagnosis of COVID–19; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the revocation of the Emergency Use Authorizations (EUAs) (the Authorizations) issued to the University of Texas MD Anderson Cancer Center, Molecular Diagnostics Laboratory (MD Anderson) for the MD Anderson High-throughput SARS–CoV–2 RT–PCR Assay, and Visby Medical, Inc. for the Visby Medical COVID–19 and Visby Medical COVID–19 Point of Care Test. FDA revoked these Authorizations under the Federal Food, Drug, and Cosmetic Act (FD&C Act). The revocations, which include an explanation of the reasons for each revocation, are reprinted in this document.

DATES: The Authorization for the MD Anderson High-throughput SARS–CoV–2 RT–PCR Assay is revoked as of November 30, 2022. The Authorizations for the Visby Medical COVID–19 and Visby Medical COVID–19 Point of Care Test are revoked as of December 2, 2022.

ADDRESSES: Submit written requests for a single copy of the revocations to the Office of Counterterrorism and Emerging Threats, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 1, Rm. 4338, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your request or include a fax number to which the revocations may be sent. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the revocations.

FOR FURTHER INFORMATION CONTACT: Jennifer Ross, Office of Counterterrorism and Emerging Threats, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 1, Rm. 4332, Silver Spring, MD 20993–0002, 301–796–8510 (this is not a toll-free number).

SUPPLEMENTARY INFORMATION:

I. Background

Section 564 of the FD&C Act (21 U.S.C. 360bbb–3) as amended by the Project BioShield Act of 2004 (Pub. L. 108–276) and the Pandemic and All-Hazards Preparedness Reauthorization Act of 2013 (Pub. L. 113–5) allows FDA to strengthen the public health protections against biological, chemical, nuclear, and radiological agents. Among other things, section 564 of the FD&C Act allows FDA to authorize the use of an unapproved medical product or an unapproved use of an approved medical product in certain situations. On June 24, 2020, FDA issued an EUA to MD Anderson for the MD Anderson High-throughput SARS–CoV–2 RT–PCR Assay, subject to the terms of the Authorization. Notice of the issuance of this Authorization was published in the **Federal Register** on November 20, 2020 (85 FR 74346), as required by section 564(h)(1) of the FD&C Act. On September 16, 2020, FDA issued an EUA to Visby Medical, Inc. for the Visby Medical COVID–19, subject to the terms of the Authorization. Notice of the issuance of this Authorization was published in the **Federal Register** on April 23, 2021 (86 FR 21749), as required by section 564(h)(1) of the FD&C Act. On February 8, 2021, FDA issued an EUA to Visby Medical, Inc. for the Visby Medical COVID–19 Point of Care Test, subject to the terms of the Authorization. Notice of the issuance of this Authorization was published in the **Federal Register** on April 23, 2021 (86 FR 21749), as required by section 564(h)(1) of the FD&C Act. Subsequent revisions to the Authorizations were made available on FDA's website. The authorization of a device for emergency use under section 564 of the FD&C Act may, pursuant to section 564(g)(2) of the FD&C Act, be revoked when the criteria under section 564(c) of the FD&C Act for issuance of such authorization are no longer met (section 564(g)(2)(B) of the FD&C Act), or other circumstances make such revocation appropriate to protect