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

the public health or safety (section 564(g)(2)(C) of the FD&C Act).

**II. EUA Revocation Requests**

On November 18, 2022, FDA received a request from MD Anderson for the withdrawal of, and on November 30, 2022, FDA revoked, the Authorization for the MD Anderson High-throughput SARS–CoV–2 RT–PCR Assay. Because MD Anderson requested FDA withdraw the EUA for the MD Anderson High-throughput SARS–CoV–2 RT–PCR Assay, FDA has determined that it is appropriate to protect the public health or safety to revoke this Authorization. On November 29, 2022, FDA received a request from Visby Medical, Inc. for the closure of, and on December 2, 2022, FDA revoked, the Authorizations for the Visby Medical COVID–19 and Visby Medical COVID–19 Point of Care Test. Because Visby Medical, Inc. requested FDA close the EUAs for the Visby Medical COVID–19 and Visby Medical COVID–19 Point of Care Test, FDA has determined that it is appropriate to protect the public health or safety to revoke these Authorizations.

**III. Electronic Access**

An electronic version of this document and the full text of the revocations are available on the internet at *https://www.regulations.gov/*.

**IV. The Revocations**

Having concluded that the criteria for revocation of the Authorizations under section 564(g)(2)(C) of the FD&C Act are met, FDA has revoked the EUAs for MD Anderson's MD Anderson High-throughput SARS–CoV–2 RT–PCR Assay and for Visby Medical, Inc.'s Visby Medical COVID–19 and Visby Medical COVID–19 Point of Care Test. The revocations in their entirety follow and provide an explanation of the reasons for each revocation, as required by section 564(h)(1) of the FD&C Act.

BILLING CODE 4164–01–P

U.S. FOOD & DRUG ADMINISTRATION

November 30, 2022

Keyur P. Patel, Ph.D.
Medical Director, Molecular Diagnostics Laboratory
The University of Texas MD Anderson Cancer Center
6565 MD Anderson Blvd.
Houston, TX 77030

**Re: Revocation of EUA200158**

Dear Dr. Patel:

This letter is in response to the request from the University of Texas MD Anderson Cancer Center, Molecular Diagnostics Laboratory, received via email on November 18, 2022, that the U.S. Food and Drug Administration (FDA) withdraw the EUA for the MD Anderson High-throughput SARS-CoV-2 RT-PCR Assay issued on June 24, 2020, and amended on September 23, 2021 and November 29, 2021. The University of Texas MD Anderson Cancer Center, Molecular Diagnostics Laboratory discontinued testing with the MD Anderson High-throughput SARS-CoV-2 RT-PCR Assay as of October 31, 2022.

The authorization of a device for emergency use under section 564 of the Federal Food, Drug, and Cosmetic Act (the Act) (21 U.S.C. 360bbb-3) may, pursuant to section 564(g)(2) of the Act, be revoked when circumstances make such revocation appropriate to protect the public health or safety (section 564(g)(2)(C) of the Act). Because the University of Texas MD Anderson Cancer Center, Molecular Diagnostics Laboratory has requested FDA withdraw the EUA for the MD Anderson High-throughput SARS-CoV-2 RT-PCR Assay, FDA has determined that it is appropriate to protect the public health or safety to revoke this authorization. Accordingly, FDA hereby revokes EUA200158 for the MD Anderson High-throughput SARS-CoV-2 RT-PCR Assay, pursuant to section 564(g)(2)(C) of the Act. As of the date of this letter, the MD Anderson High-throughput SARS-CoV-2 RT-PCR Assay is no longer authorized for emergency use by FDA.

Notice of this revocation will be published in the *Federal Register*, pursuant to section 564(h)(1) of the Act.

Sincerely,

/s/

Namandjé N. Bumpus, Ph.D.
Chief Scientist
Food and Drug Administration

U.S. FOOD & DRUG ADMINISTRATION

December 2, 2022

Beth Lingenfelter
VP of Clinical and Regulatory Affairs
Visby Medical, Inc.
3010 North First Street
San Jose, CA 95134

**Re: Revocation of EUA202677**

Dear Ms. Lingenfelter:

This letter is in response to the request from Visby Medical, Inc., received via email on November 29, 2022, that the U.S. Food and Drug Administration (FDA) close the EUA for the Visby Medical COVID-19 issued on September 16, 2020, amended on December 28, 2020, and reissued on August 31, 2021. Visby Medical, Inc. discontinued manufacturing the Visby Medical COVID-19 test on January 25, 2021, and FDA understands there are no viable (non-expired) Visby Medical COVID-19 tests remaining in distribution in the United States.

The authorization of a device for emergency use under section 564 of the Federal Food, Drug, and Cosmetic Act (the Act) (21 U.S.C. 360bbb-3) may, pursuant to section 564(g)(2) of the Act, be revoked when circumstances make such revocation appropriate to protect the public health or safety (section 564(g)(2)(C) of the Act). Because Visby Medical, Inc. has requested FDA close the EUA for the Visby Medical COVID-19, FDA has determined that it is appropriate to protect the public health or safety to revoke this authorization. Accordingly, FDA hereby revokes EUA202677 for the Visby Medical COVID-19, pursuant to section 564(g)(2)(C) of the Act. As of the date of this letter, the Visby Medical COVID-19 is no longer authorized for emergency use by FDA.

Notice of this revocation will be published in the *Federal Register*, pursuant to section 564(h)(1) of the Act.

Sincerely,

/s/

Namandjé N. Bumpus, Ph.D.
Chief Scientist
Food and Drug Administration

U.S. FOOD & DRUG ADMINISTRATION

December 2, 2022

Beth Lingenfelter
VP of Clinical and Regulatory Affairs
Visby Medical, Inc.
3010 North First Street
San Jose, CA 95134

**Re: Revocation of EUA203089**

Dear Ms. Lingenfelter:

This letter is in response to the request from Visby Medical, Inc., received via email on November 29, 2022, that the U.S. Food and Drug Administration (FDA) close the EUA for the Visby Medical COVID-19 Point of Care Test issued on February 8, 2021, amended on September 23, 2021, and reissued on July 22, 2022. Visby Medical, Inc. discontinued manufacturing the Visby Medical COVID-19 Point of Care Test on April 25, 2022, and FDA understands there are no viable (non-expired) Visby Medical COVID-19 Point of Care Tests remaining in distribution in the United States.

The authorization of a device for emergency use under section 564 of the Federal Food, Drug, and Cosmetic Act (the Act) (21 U.S.C. 360bbb-3) may, pursuant to section 564(g)(2) of the Act, be revoked when circumstances make such revocation appropriate to protect the public health or safety (section 564(g)(2)(C) of the Act). Because Visby Medical, Inc. has requested FDA close the EUA for the Visby Medical COVID-19 Point of Care Test, FDA has determined that it is appropriate to protect the public health or safety to revoke this authorization. Accordingly, FDA hereby revokes EUA203089 for the Visby Medical COVID-19 Point of Care Test, pursuant to section 564(g)(2)(C) of the Act. As of the date of this letter, the Visby Medical COVID-19 Point of Care Test is no longer authorized for emergency use by FDA.

Notice of this revocation will be published in the *Federal Register*, pursuant to section 564(h)(1) of the Act.

Sincerely,

/s/

Namandjé N. Bumpus, Ph.D.
Chief Scientist
Food and Drug Administration

Dated: December 27, 2022.

**Lauren K. Roth,**

*Associate Commissioner for Policy.*

[FR Doc. 2022–28496 Filed 12–30–22; 8:45 am]

**BILLING CODE 4164–01–C**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Announcement of First Meeting of the 2025 Dietary Guidelines Advisory Committee and Request for Comments

**AGENCY:** U.S. Department of Health and Human Services (HHS), Office of the Assistant Secretary for Health (OASH); U.S. Department of Agriculture (USDA), Food, Nutrition, and Consumer Services (FNCS).

**ACTION:** Notice.

**SUMMARY:** The U.S. Departments of Health and Human Services and Agriculture announce the first meeting of the newly appointed 2025 Dietary Guidelines Advisory Committee (Committee). This meeting will be open to the public virtually. Additionally, this notice opens a public comment period that will remain open until late 2024, throughout the Committee's deliberations.

**DATES:** The first meeting will be held February 9–10, 2023. The public comment period opens with the publication of this notice.

**ADDRESSES:**

(a) The meeting will be accessible online via livestream and recorded for later viewing. Registrants will receive the livestream information prior to the meeting.

(b) You may send comments, identified by Docket OASH–2022–0021, by either of the following methods:

- *Online (preferred method):* Federal eRulemaking Portal: *http://www.regulations.gov.*
- *Mail:* Janet M. de Jesus, MS, RD, HHS/OASH Office of Disease Prevention and Health Promotion