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 Government Accountability Office (GAO) is now accepting nominations for MACPAC appointments that will be effective May 2025. 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 28, 2025, 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: Corissa Kiyan-Fukumoto at *KiyanFukumotoC@gao.gov* or (202) 512–7114 if you do not receive an acknowledgment or need additional information. For general information, contact GAO's Office of Public Affairs, at *PublicAffairs@gao.gov*.

Authority: 42 U.S.C. 1396.

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2024–29982 Filed 12–31–24; 8:45 am]

BILLING CODE 1610–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of Meeting; National Advisory Committee on the Trafficking of Children and Youth in the United States

AGENCY: Office on Trafficking in Persons, Administration for Children and Families, Department of Health and Human Services.

ACTION: Announcement of meeting.

SUMMARY: Notice is hereby given, pursuant to the provisions of the Federal Advisory Committee Act (FACA), that a meeting of the National Advisory Committee on the Trafficking of Children and Youth in the United States (Committee) will be held on January 13, 2025. The purpose of the meeting is for the Committee to discuss its plans and responsibilities over its 24-month extension to advise on the development and implementation of successful interventions with children and youth who have been impacted by labor and sex trafficking, make recommendations for administrative and legislative changes regarding the trafficking of children and youth, and publish information on best practices regarding the labor and sex trafficking of children and youth.

DATES: Please submit any comments regarding best practices and recommendations to address the trafficking of children and youth comments as soon as possible and before January 10, 2025. The meeting will be held on January 13, 2025. Future meetings will be held on a quarterly basis, on or around the approximate dates: April 14, 2025; July 14, 2025; and October 13, 2025.

ADDRESSES: The meeting will be held virtually. Please register for this event online: *https://www.acf.hhs.gov/otip/partnerships/nac-trafficking-children-youth*. Please submit comments to *endtrafficking@acf.hhs.gov* with the subject ''NAC Comments''.

FOR FURTHER INFORMATION CONTACT: Katherine Chon (Designated Federal Officer) at *EndTrafficking@acf.hhs.gov* or (202) 205–5778, or 330 C Street SW, Washington, DC 20201. Additional information is available at *https://www.acf.hhs.gov/otip/partnerships/nac-trafficking-children-youth*.

SUPPLEMENTARY INFORMATION: The formation and operation on behalf of the Committee are governed by the provisions of Public Law 92–463, as amended (5 U.S.C. App. 2), which sets forth standards for the formation and use of Federal advisory committees.

Purpose of the Committee: The purpose of the Committee is to advise the Secretary and the Attorney General on practical and general policies concerning improvements to the Nation's response to the labor and sex trafficking of children and youth in the United States. The discretionary advisory Committee will build on the work of the previous National Advisory Committee on the Sex Trafficking of Children and Youth in the United States, which operated from January 18, 2017, to January 18, 2022, authorized pursuant to section 121 of the Preventing Sex Trafficking and Strengthening Families Act (Pub. L. 113–183) and governed by the provisions of Public Law 92–463, as amended 5 U.S.C. chapter 10.

Tentative Agenda: The agenda can be found at *https://www.acf.hhs.gov/otip/partnerships/nac-trafficking-children-youth.* To submit written statements, email *endtrafficking@acf.hhs.gov* by January 10, 2025. Please include your name, organization, and phone number. More details on these options are below.

Public Accessibility to the Meeting: Pursuant to 5 U.S.C. 552b and 41 CFR 102–3.140 through 102–3.165, and subject to the availability of space, this meeting is open to the public virtually.

Written Statements: Pursuant to 41 CFR 102–3.105(j) and 102–3.140 and section 10(a)(3) of the Federal Advisory Committee Act, the public may submit written statements in response to the stated agenda of the meeting or to the Committee's mission in general. Organizations with recommendations on strategies to engage states and stakeholders are encouraged to submit their comments or resources (hyperlinks preferred). Written comments or statements received after January 10, 2025, may not be provided to the Committee until its next meeting.

Verbal Statements: Pursuant to 41 CFR 102–3.140, the Committee is not obligated to allow a member of the public to speak or otherwise address the Committee during the meeting. Members of the public are invited to provide verbal statements during the Committee meeting only at the time and manner described in the agenda. The request to speak should include a brief statement of the subject matter to be addressed and should be relevant to the stated agenda of the meeting or the Committee's mission in general.

Minutes: The minutes of this meeting will be available for public review and copying within 90 days at *https://www.acf.hhs.gov/otip/partnerships/nac-trafficking-children-youth.*

Rudette Pinkney,

Acting Deputy Director, Office of the Executive Secretariat.

[FR Doc. 2024–31416 Filed 12–31–24; 8:45 am]

BILLING CODE 4184–40–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–1991–P–0355]

Liquid Eggs Deviating From the Standard of Identity; Revocation of Temporary Permit for Market Testing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the revocation of the temporary permit issued to M.G. Waldbaum Co., a subsidiary of Michael Foods Egg Co., to market test ''ultrapasteurized liquid whole eggs'' and ''ultrapasteurized liquid whole eggs with citric acid'' because the need for the temporary permit no longer exists.

DATES: This permit is revoked as of January 2, 2025.

FOR FURTHER INFORMATION CONTACT: Marjan Morravej, Nutrition Center of Excellence, Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–2371; or Jessica Ritsick, Office of Policy, Regulations, and Information, Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–2378.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 21, 1989 (54 FR 30612), we issued a notice announcing that we had issued a temporary permit to Crystal Foods, Inc., 6465 Wayzata Blvd., Minneapolis, MN 55426, a subsidiary of Michael Foods, Inc., to market test experimental packs of ''ultrapasteurized liquid whole eggs'' and ''ultrapasteurized liquid whole eggs with citric acid,'' which we stated deviate from the standard of identity for liquid eggs at § 160.115 (21 CFR 160.115) because they were processed with increased heat treatment and aseptic processing and packaging. We refer to the temporary permit holder as ''the company'' throughout this notice. The temporary permit allowed the company to measure consumer acceptance of the product, identify mass production problems, and assess commercial feasibility (Id.). In February 1991, FDA combined the original docket for the temporary permit (FDA–1989–P–0168) with other related dockets for the company into what is now docket number FDA–1991–P–0355.

After issuance of the temporary permit, the company requested, and FDA granted, several revisions:

- July 11, 1990 (55 FR 28456)—FDA amended the temporary permit to provide for package sizes larger than the designated 2.27 kilograms (5 pounds) to provide a broader base for data collection on consumer acceptance of the test products.
- September 20, 1990 (55 FR 38753)—FDA extended the temporary permit so the company could continue experimental market testing of the products and continue gathering data in support of its petition to amend the standard of identity for liquid eggs at § 160.115. As part of the extension, FDA invited interested persons to participate in the market test under the conditions in the temporary permit, except for the designated area of distribution. We have no records that show that any interested persons notified us of their intent to participate in the market test, as required under § 130.17(i) (21 CFR 130.17(i)).
- March 22, 1991 (56 FR 12206)—FDA amended the temporary permit to allow the test products to be packaged in aseptic packages ranging in size from 42.5 grams (1.5 ounces) to 1 kilogram (2.2 pounds). Additionally, as requested by the company, we changed the name and address of the permit holder from Crystal Foods, Inc., Minneapolis, MN 55426, to M.G. Waldbaum Co., Wakefield, NE 68784.

In the time since the temporary permit was originally issued, FDA has concluded that the temporary permit is not necessary, because the standard of identity in § 160.115 provides for the treatment process used by the company under the temporary permit. Our regulation, at § 160.115(a), states that liquid eggs must be pasteurized or otherwise treated to destroy all viable *Salmonella* microorganisms. The specific process used by the company under the temporary permit—increased heat treatment and aseptic processing and packaging—is consistent with § 160.115(a). Specifically, the standard of identity for liquid eggs permits other treatments that destroy all viable *Salmonella* microorganisms. As such, we have concluded that the temporary permit is not necessary to market liquid eggs using the company's process, consistent with the standard of identity.

In addition, in April 2024, FDA contacted the company via email regarding the current use of its temporary permit. The company did not object to FDA revoking the temporary permit under § 130.17(g)(3).

Therefore, under § 130.17(g)(3), we are revoking the company's temporary permit because the need for it no longer exists.

Dated: December 20, 2024.

P. Ritu Nalubola,

Associate Commissioner for Policy.

[FR Doc. 2024–31470 Filed 12–31–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–D–4974]

Advanced Manufacturing Technologies Designation Program; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance for industry entitled ''Advanced Manufacturing Technologies Designation Program.'' FDA encourages the early adoption of advanced manufacturing technologies (AMTs) by the pharmaceutical industry, which can improve the reliability and robustness of the manufacturing process and can benefit patients by enhancing product quality and reducing drug development time or increasing or maintaining the supply of drugs that are life-supporting, life-sustaining, of critical importance to providing health care, or in shortage. This guidance provides recommendations to persons and organizations interested in participating in FDA's Advanced Manufacturing Technologies Designation Program, which facilitates the development of drugs manufactured using an AMT that has been designated as such under the program. The guidance finalizes the draft guidance of the same title issued on December 13, 2023.

DATES: The announcement of the guidance is published in the **Federal Register** on January 2, 2025.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment 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