Records generated from these meetings may be inspected and reproduced at the Regional Programs Coordination Unit Office, as they become available, both before and after each meeting. Records of the meetings will be available via the file sharing website, *https://bit.ly/4g3IB4K.* Persons interested in the work of this Committee are directed to the Commission's website, *http://www.usccr.gov,* or may contact the Regional Programs Coordination Unit at *ebohor@usccr.gov.*

Agenda

I. Welcome and Roll Call
II. Approval of Minutes
III. Announcements and Updates
IV. Discussion
  a. Project Proposal
  b. Begin planning for public briefings on antisemitism in Ohio
V. Next steps
VI. Public Comment
VII. Adjournment

Dated: March 11, 2025.

David Mussatt,

Supervisory Chief, Regional Programs Unit.

[FR Doc. 2025–04138 Filed 3–14–25; 8:45 am]

BILLING CODE P

COMMISSION ON CIVIL RIGHTS

Notice of Public Meeting of the District of Columbia Advisory Committee to the U.S. Commission on Civil Rights

AGENCY: U.S. Commission on Civil Rights.

ACTION: Notice of virtual business meeting.

SUMMARY: Notice is hereby given, pursuant to the provisions of the rules and regulations of the U.S. Commission on Civil Rights (Commission) and the Federal Advisory Committee Act, that the District of Columbia Advisory Committee (Committee) to the U.S. Commission on Civil Rights will hold a public meeting via Zoom. The purpose of the meeting is to discuss continued activities on the committee's topic of accessibility and provision of special education for students with disabilities in DC public schools.

DATES: Thursday, April 3, 2025, from 12 p.m.–1 p.m. Eastern Time.

ADDRESSES: The meeting will be held via Zoom.

Registration Link (Audio/Visual): https://tinyurl.com/2pt5mvdw.

Join by Phone (Audio Only): 1–833–435–1820 USA Toll Free; Webinar ID: 161 197 6779#.

FOR FURTHER INFORMATION CONTACT: Melissa Wojnaroski, DFO, at *mwojnaroski@usccr.gov* or 1–202–618–4158.

SUPPLEMENTARY INFORMATION: This Committee meeting is available to the public through the registration link above. Any interested member of the public may attend this meeting. An open comment period will be provided to allow members of the public to make oral statements as time allows. Pursuant to the Federal Advisory Committee Act, public minutes of the meeting will include a list of persons who are present at the meeting. If joining via phone, callers can expect to incur regular charges for calls they initiate over wireless lines, according to their wireless plan. The Commission will not refund any incurred charges. Callers will incur no charge for calls they initiate over land-line connections to the toll-free telephone number. Closed captioning is available by selecting ''CC'' in the meeting platform. To request additional accommodations, please email *ebohor@usccr.gov* at least 10 business days prior to the meeting.

Members of the public are entitled to submit written comments; the comments must be received in the regional office within 30 days following the scheduled meeting. Written comments may be emailed to Evelyn Bohor at *ebohor@usccr.gov.* Persons who desire additional information may contact the Regional Programs Coordination Unit at 1–202–809–9618.

Records generated from this meeting may be inspected and reproduced at the Regional Programs Coordination Unit Office, as they become available, both before and after the meeting. Records of the meetings will be available via *www.facadatabase.gov* under the Commission on Civil Rights, District of Columbia Advisory Committee link. Persons interested in the work of this Committee are directed to the Commission's website, *http://www.usccr.gov,* or may contact the Regional Programs Coordination Unit at *ebohor@usccr.gov.*

Agenda

I. Welcome and Roll Call
II. Approval of Minutes
III. Discussion
IV. Public Comment
V. Next Steps
VI. Adjournment

Dated: March 11, 2025.

David Mussatt,

Supervisory Chief, Regional Programs Unit.

[FR Doc. 2025–04137 Filed 3–14–25; 8:45 am]

BILLING CODE 6335–01–P

DEPARTMENT OF COMMERCE

Foreign-Trade Zones Board

[B–15–2025]

Foreign-Trade Zone (FTZ) 43, Notification of Proposed Production Activity; Pfizer, Inc.; (Pharmaceutical Intermediate Product); Kalamazoo, Michigan

Pfizer, Inc. (Pfizer) submitted a notification of proposed production activity to the FTZ Board (the Board) for its facilities in Kalamazoo, Michigan within Subzone 43E. The notification conforming to the requirements of the Board's regulations (15 CFR 400.22) was received on March 10, 2025.

Pursuant to 15 CFR 400.14(b), FTZ production activity would be limited to the specific foreign-status material(s)/component(s) and specific finished product(s) described in the submitted notification (summarized below) and subsequently authorized by the Board. The benefits that may stem from conducting production activity under FTZ procedures are explained in the background section of the Board's website—accessible via *www.trade.gov/ftz.* The proposed finished product and material/component would be added to the production authority that the Board previously approved for the operation, as reflected on the Board's website.

The proposed finished product is phthalimido lactol (pharmaceutical intermediate product) (duty rate—6.5%).

The proposed foreign-status material/component is 3 phthalimidopropionaldehyde (3-PPA) (duty rate—6.5%). The request indicates that 3-PPA is subject to duties under section 1702(a)(1)(B) of the International Emergency Economic Powers Act (section 1702) and section 301 of the Trade Act of 1974 (section 301), depending on the country of origin. The applicable section 1702 and section 301 decisions require subject merchandise to be admitted to FTZs in privileged foreign status (19 CFR 146.41).

Public comment is invited from interested parties. Submissions shall be addressed to the Board's Executive Secretary and sent to: *ftz@trade.gov.* The closing period for their receipt is April 28, 2025.

A copy of the notification will be available for public inspection in the ''Online FTZ Information System'' section of the Board's website.

For further information, contact Diane Finver at *Diane.Finver@trade.gov.*

Dated: March 11, 2025.

Elizabeth Whiteman,

Executive Secretary.

[FR Doc. 2025–04257 Filed 3–14–25; 8:45 am]

BILLING CODE 3510–DS–P

DEPARTMENT OF COMMERCE

International Trade Administration

[A–570–992]

Monosodium Glutamate From the People's Republic of China: Preliminary Affirmative Determination of Circumvention; Correction

AGENCY: Enforcement and Compliance, International Trade Administration, Department of Commerce.

ACTION: Notice; correction.

SUMMARY: The U.S. Department of Commerce (Commerce) published notice in the **Federal Register** of February 21, 2025 of its preliminary affirmative determination of circumvention of the antidumping duty order on monosodium glutamate (MSG) from the People's Republic of China (China). In that notice, the importer and exporter certifications provided at Appendix II contained three errors.

FOR FURTHER INFORMATION CONTACT: Thomas Cloyd, AD/CVD Operations, Office VII, Enforcement and Compliance, International Trade Administration, U.S. Department of Commerce, 1401 Constitution Avenue NW, Washington, DC 20230; telephone: (202) 482–1246.

SUPPLEMENTARY INFORMATION:

Background

On February 21, 2025, Commerce published in the **Federal Register** its *Preliminary Determination* of circumvention of the antidumping duty order on MSG from China.[1] In the *Preliminary Determination,* the importer and exporter certifications provided at Appendix II contained three errors. Commerce is hereby correcting the certifications at Appendix II to be in accordance with its preliminary decision. The corrected Appendix II is attached to this notice.

Correction

First, in the **Federal Register** of February 21, 2025, in FR Doc 2025–02924 on page 10071, in the second column, correct Appendix II by replacing the entire text of section (J) of the importer certification as follows:

''I understand that {IMPORTING COMPANY} is required to submit a copy of the importer and exporter certifications as part of the entry summary by uploading them into the document imaging system (DIS) in ACE, and to provide U.S. Customs and Border Protection (CBP) and/or the U.S. Department of Commerce (Commerce) with the importer certification, and any supporting documentation, and a copy of the exporter's certification, and any supporting documentation provided to the importer by the exporter, upon request of either agency.''

Second, in the **Federal Register** of February 21, 2025, in FR Doc 2025–02924 on page 10071, in the third column, correct Appendix II by replacing the entire text of section (M) of the importer certification as follows:

''I understand that agents of the importer, such as brokers, are not permitted to make this certification. This certification was completed by the time of filing the entry summary or within 45 days of the date on which Commerce published notice of its preliminary circumvention findings in the **Federal Register**.''

Third, in the **Federal Register** of February 21, 2025, in FR Doc 2025–02924 on page 10072, in the first and second columns, correct Appendix II by replacing the entire text of section (L) of the exporter certification, as follows:

''This certification was completed at time of shipment or within 45 days of the date on which Commerce published notice of its preliminary circumvention findings in the **Federal Register**.''

Notification to Interested Parties

This notice is issued and published in accordance with section 781(b) of the Tariff Act of 1930, as amended, and 19 CFR 351.226.

Dated: March 12, 2025.

Christopher Abbott,

Deputy Assistant Secretary for Policy and Negotiations, performing the non-exclusive functions and duties of the Assistant Secretary for Enforcement and Compliance.

Appendix II—Importer Certification

I hereby certify that:

(A) My name is {IMPORTING COMPANY OFFICIAL'S NAME} and I am an official of {NAME OF IMPORTING COMPANY}, located at {ADDRESS OF IMPORTING COMPANY}.

(B) I have direct personal knowledge of the facts regarding the importation into the Customs territory of the United States of the monosodium glutamate (MSG) assembled or completed in Malaysia that entered under the entry summary number(s), identified below, and are covered by this certification. ''Direct personal knowledge'' refers to facts the certifying party is expected to have in its own records. For example, the importer must have direct personal knowledge of the importation of the product, including the exporter's and/or foreign seller's identity and location.

(C) If the importer is acting on behalf of the first U.S. customer, include the following sentence as paragraph C of this certification:

The MSG covered by this certification was imported by {IMPORTING COMPANY} on behalf of {U.S. CUSTOMER}, located at {ADDRESS OF U.S. CUSTOMER}.

If the importer is not acting on behalf of the first U.S. customer, include the following sentence as paragraph C of this certification:

{NAME OF IMPORTING COMPANY} is not acting on behalf of the first U.S. customer.

(D) The MSG covered by this certification was shipped to {NAME OF PARTY IN THE UNITED STATES TO WHOM THE MERCHANDISE WAS FIRST SHIPPED}, located at {U.S. ADDRESS TO WHICH MERCHANDISE WAS SHIPPED}.

(E) I have personal knowledge of the facts regarding the production of the imported products covered by this certification. ''Personal knowledge'' includes facts obtained from another party, (*e.g.,* correspondence received by the importer (or exporter) from the producer regarding the source of the inputs (*i.e.,* glutamic acid) used to produce the imported MSG).

(F) This certification applies to the following entries (repeat this block as many times as necessary):

Entry Summary #:
Entry Summary Line Item #:
Foreign Seller:
Foreign Seller's Address:
Foreign Seller's Invoice #:
Foreign Seller's Invoice Line Item #:
Country of Origin of Glutamic Acid:
Producer:
Producer's Address:

(G) The MSG covered by this certification was not produced using glutamic acid produced in the People's Republic of China.

(H) I understand that {IMPORTING COMPANY} is required to maintain a copy of this certification and sufficient documentation supporting this certification (*i.e.,* documents maintained in the normal course of business, or documents obtained by the certifying party, for example, certificates of origin, product data sheets, mill test reports, productions records, invoices, etc.) until the later of: (1) the date that is five years after the date of the latest entry covered by the certification; or (2) the date that is three years after the conclusion of any litigation in the United States courts regarding such entries.

(I) I understand that {IMPORTING COMPANY} is required to maintain a copy of the exporter's certification (attesting to the production and/or exportation of the imported merchandise identified above), and any supporting documentation provided to the importer by the exporter, until the later of: (1) the date that is five years after the date of the latest entry covered by the certification; or (2) the date that is three years after the conclusion of any litigation in United States courts regarding such entries.

(J) I understand that {IMPORTING COMPANY} is required to submit a copy of

[1] *See Monosodium Glutamate from the People's Republic of China: Preliminary Affirmative Determination of Circumvention,* 90 FR 10068 (February 21, 2025) (*Preliminary Determination*); *see also Monosodium Glutamate from the People's Republic of China: Second Amended Final Determination of Sales at Less Than Fair Value and Amended Antidumping Order,* 80 FR 487 (January 6, 2015) (*Order*).