be accessed through the following link Webex: https://bit.ly/NYSC51921.

FOR FURTHER INFORMATION CONTACT: Ana Victoria Fortes, Designated Federal Officer (DFO) at afortes@usccr.gov or by phone at (202) 681–0857.

SUPPLEMENTARY INFORMATION: This meeting is available to the public through the following toll-free call-in number: 800–360–9505; Access code: 199 070 2234. Any interested member of the public may call this number and listen to the meeting. Callers can expect to incur charges for calls they initiate over wireless lines, and 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.

Persons with hearing impairments may also follow the proceedings by first calling the Federal Relay Service at 1–800–877–8339 and providing the Service with the conference call number and conference ID number.

Members of the public are entitled to make comments during the open period at the end of the meeting. Members of the public may also submit written comments; the comments must be received in the Regional Programs Unit within 30 days following the meeting. Written comments may be mailed to the Western Regional Office, U.S. Commission on Civil Rights, 300 North Los Angeles Street, Suite 2010, Los Angeles, CA 90012 or email Ana Victoria Fortes at afortes@usccr.gov.

Records and documents discussed during the meeting will be available for public viewing prior to and after the meeting at https://www.facadatabase.gov/FACA/FACAPublicViewCommitteeDetails?id=a1000000011gjzJAAQ.

Please click on the “Committee Meetings” tab. Records generated from these meetings may also be inspected and reproduced at the Regional Programs Unit, as they become available, both before and after the meetings. Persons interested in the work of this Committee are directed to the Commission’s website, https://www.usccr.gov, or may contact the Regional Programs Unit at the above email or street address.

Agenda

Opening Remarks (12:00–12:15 p.m.)
   - Erin Phillips, President,
     Power2Parent
   - Representative, Nevada COVID–19
     Response, Relief and Recovery
     Taskforce
   - Rebecca Garcia, President, Nevada
     Parent Teacher Association
Q & A (1:15–2:10 p.m.)
Public Comment (2:10–2:25 p.m.)
Closing Remarks (2:25–2:30 p.m.)

Dated: March 11, 2021.
David Mussatt,
Supervisory Chief, Regional Programs Unit.
[FR Doc. 2021–05471 Filed 3–16–21; 8:45 am]
BILLING CODE P

DEPARTMENT OF COMMERCE

Foreign Trade Zones Board
[B–20–2021]

Foreign Trade Zone (FTZ) 177—Evansville, Indiana; Notification of Proposed Production Activity; AstraZeneca Pharmaceuticals LP (Pharmaceutical Products); Mount Vernon, Indiana

AstraZeneca Pharmaceuticals LP (AstraZeneca) submitted a notification of proposed production activity to the FTZ Board for its facility in Mount Vernon, Indiana. The notification conforming to the requirements of the regulations of the FTZ Board (15 CFR 400.22) was received on March 9, 2021. AstraZeneca already has authority to produce certain pharmaceutical products within Subzone 177A. The current request would add finished products and foreign status materials to the scope of authority. Pursuant to 15 CFR 400.14(b), additional FTZ authority would be limited to the specific foreign-status materials and specific finished products described in the submitted notification (as described below) and subsequently authorized by the FTZ Board.

Production under FTZ procedures could exempt AstraZeneca from customs duty payments on the foreign-status materials/components used in export production. On its domestic sales, for the foreign-status materials/components noted below and in the existing scope of authority. AstraZeneca would be able to choose the duty rates during customs entry procedures that apply to: CALQUENCE (acalabrutinib) capsules; DAKLINZA (daclatasvir) tablets; FARXIGA/FORXIGA (dapagliflozin) tablets; KOMBIGLYZE IR (metformin hydrochloride and saxagliptin hydrochloride) tablets; KOMBIGLYZE XR (metformin hydrochloride and saxagliptin hydrochloride) tablets; METFORMIN IR (metformin hydrochloride) tablets; OGLYZA (saxagliptin hydrochloride) tablets; QTERN (dapagliflozin and saxagliptin hydrochloride) tablets; QTERNMET XR (dapagliflozin, metformin hydrochloride and saxagliptin hydrochloride) tablets; TARGSISSO (osimertinib mesylate) tablets; XIGDUO IR (dapagliflozin and metformin hydrochloride) tablets; and, XIGDUO XR (dapagliflozin and metformin hydrochloride) tablets (duty-free). AstraZeneca would be able to avoid duty on foreign-status components which become scrap/waste. Customs duties also could possibly be deferred or reduced on foreign-status production equipment.

The materials sourced from abroad include: Metformin hydrochloride active pharmaceutical ingredient (API); dapagliflozin API; daclatasvir API; osimertinib mesylate API; acalabrutinib API; and, saxagliptin hydrochloride API (duty rate ranges from 3.7% to 6.5%). The request indicates that certain materials are subject to duties under Section 301 of the Trade Act of 1974 (Section 301), depending on the country of origin. The applicable 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 26, 2021.

A copy of the notification will be available for public inspection in the “Reading Room” section of the Board’s website, which is accessible via www.trade.gov/ftz.

For further information, contact Christopher Wedderburn at Chris.Wedderburn@trade.gov.

Dated: March 11, 2021.
Andrew McGilvray,
Executive Secretary.
[FR Doc. 2021–05478 Filed 3–16–21; 8:45 am]
BILLING CODE 3510–DS–P

DEPARTMENT OF COMMERCE

International Trade Administration

[C–531–846]

Certain Hot-Rolled Steel Flat Products From Brazil: Rescission of the 2019 Countervailing Duty Administrative Review

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

SUMMARY: The Department of Commerce (Commerce) is rescinding the administrative review of the countervailing duty (CVD) order on certain hot-rolled steel flat products (hot-rolled steel) from Brazil for the
period of review (POR) January 1, 2019, through December 31, 2019.


Background

On October 1, 2020, Commerce published in the Federal Register a notice of opportunity to request an administrative review of the CVD order on hot-rolled steel from Brazil for the POR.1 On October 30, 2020, Commerce received a timely request from AK Steel Corporation, Nucor Corporation, United States Steel Corporation, Steel Dynamics, Inc., and SSAB Enterprises, LLC (collectively, domestic interested parties), in accordance with section 751(a) of the Tariff Act of 1930, as amended (the Act), and 19 CFR 351.213(b), to conduct an administrative review of this CVD order for 12 companies.2

On December 8, 2020, Commerce published in the Federal Register a notice of initiation with respect to these companies.3 On February 2, 2021, the domestic interested parties timely withdrew their request for an administrative review for all 12 companies.4

Rescission of Review

Pursuant to 19 CFR 351.213(d)(1), Commerce will rescind an administrative review, in whole or in part, if the parties that requested a review withdraw the request within 90 days of the date of publication of notice of initiation of the requested review. As noted above, the domestic interested parties withdrew their request for review by the 90-day deadline, and no other party requested an administrative review of this order. Therefore, we are rescinding the administrative review of the CVD order on certain hot-rolled steel flat products from Brazil covering the period January 1, 2019, through December 31, 2019, in its entirety.

Assessment

Commerce will instruct U.S. Customs and Border Protection (CBP) to assess countervailing duties on all appropriate entries. Because Commerce is rescinding this administrative review in its entirety, the entries to which this administrative review pertained shall be assessed at rates equal to the cash deposit of estimated countervailing duties required at the time of entry, or withdrawal from warehouse, for consumption, in accordance with 19 CFR 351.212(c)(1)(i). Commerce intends to issue appropriate assessment instructions directly to CBP no earlier than 35 days after the date of publication of this notice in the Federal Register.

Notification Regarding Administrative Protective Orders

This notice serves as the only reminder to parties subject to administrative protective order (APO) of their responsibility concerning the return or destruction of proprietary information disclosed under APO in accordance with 19 CFR 351.305(a)(3). Timely written notification of the return/destruction of APO materials or conversion to judicial protective order is hereby requested. Failure to comply with the regulations and terms of an APO is a sanctionable violation.

Notification to Interested Parties

This notice is issued and published in accordance with sections 751(a)(1) and 777(i)(1) of the Act and 19 CFR 351.213(d)(4).

Dated: March 11, 2021.

James Maeder,
Deputy Assistant Secretary for Antidumping and Countervailing Duty Operations.

[FR Doc. 2021–05477 Filed 3–16–21; 8:45 am]
BILLING CODE 3510–05–P

DEPARTMENT OF COMMERCE
National Oceanic and Atmospheric Administration

[TID 0648–XA199]

Takes of Marine Mammals Incidental to Specified Activities; Taking Marine Mammals Incidental to Army Corps of Engineers Port San Luis Breakwater Repair Project, Avila Beach, California

AGENCY: National Marine Fisheries Service (NMFS), National Oceanic and Atmospheric Administration (NOAA), Commerce.

ACTION: Notice; proposed incidental harassment authorization; request for comments on proposed authorization and possible renewal.

SUMMARY: NMFS has received a request from the Army Corps of Engineers (ACOE) for authorization to take marine mammals incidental to the Port San Luis Breakwater Repair Project in Avila Beach, California. Pursuant to the Marine Mammal Protection Act (MMPA), NMFS is requesting comments on its proposal to issue an incidental harassment authorization (IHA) to incidentally take marine mammals during the specified activities. NMFS is also requesting comments on a possible one-year renewal that could be issued under certain circumstances and if all requirements are met, as described in the Request for Public Comments at the end of this notice. NMFS will consider public comments prior to making any final decision on the issuance of the requested MMPA authorizations and agency responses will be summarized in the final notice of our decision.

DATES: Comments and information must be received no later than April 16, 2021.

ADDRESSES: Comments should be addressed to Jolie Harrison, Chief, Permits and Conservation Division, Office of Protected Resources, National Marine Fisheries Service. Electronic comments should be sent to ITP.Meadows@noaa.gov.

Instructions: NMFS is not responsible for comments sent by any other method, to any other address or individual, or received after the end of the comment period. Comments received electronically, including all attachments, must not exceed a 25-megabyte file size. Attachments to electronic comments will be accepted in Microsoft Word or Excel or Adobe PDF file formats only. All comments received are a part of the public record and will generally be posted online at https://www.fisheries.noaa.gov/permit/incidental-take-authorizations-under-marine-mammal-protection-act without change. All personal identifying information (e.g., name, address) voluntarily submitted by the commenter may be publicly accessible. Do not submit confidential business information or otherwise sensitive or protected information.

FOR FURTHER INFORMATION CONTACT: Dwayne Meadows, Ph.D., Office of Protected Resources, NMFS, (301) 427–8401. Electronic copies of the application and supporting documents, as well as a list of the references cited in this document, may be obtained online at: https://