Proposed Rules

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This section of the FEDERAL REGISTER contains notices to the public of the proposed issuance of rules and regulations. The purpose of these notices is to give interested persons an opportunity to participate in the rule making prior to the adoption of the final rules.

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

DEPARTMENT OF THE TREASURY

19 CFR Part 190

[USCBP-2018-0029]

RIN 1515-AE23

Modernized Drawback; Correction

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security; Department of the Treasury. **ACTION:** Notice of proposed rulemaking;

correction.

SUMMARY: This document corrects a proposed regulation in a notice of proposed rulemaking published in the Federal Register of August 2, 2018, regarding Modernized Drawback. Specifically, CBP inadvertently proposed in 19 CFR 190.32(d)(2) an exemption for drawback claims for wine which included an imprecise reference to the entirety of paragraph (b). The reference should have been only to paragraphs (b)(1) and (b)(2), the specific paragraphs regarding the "lesser of" rule. As is evident from the entirety of the proposed rule, paragraph (b)(3), which implements the statutory prohibition on double drawback, applies to all drawback claims for wine. This technical correction remedies a clerical error that occurred when the language of paragraph (b)(3) was moved from a different part of the proposed regulations.

DATES: August 20, 2018.

FOR FURTHER INFORMATION CONTACT:

Randy Mitchell, U.S. Customs and Border Protection, Office of Trade, Trade Policy and Programs, 202–863– 6532, randy.mitchell@cbp.dhs.gov.

SUPPLEMENTARY INFORMATION: In proposed rule FR Doc. 2018–16279 appearing on page 37886 in the **Federal Register** issue of August 2, 2018, the following corrections are made:

1. On page 37936 in the first column, correct § 190.32 by revising paragraph (d)(2) to read as follows: § 190.32 Substitution unused merchandise drawback.

(d) * * *

(2) Allowable refund. For any drawback claim for wine (as defined in § 190.2) based on subsection (j)(2), the total amount of drawback allowable will be equal to 99 percent of the duties, taxes, and fees paid with respect to the imported merchandise, without regard to the limitations in paragraph (b)(1) or

Dated: August 14, 2018.

Robert E. Perez,

(b)(2).

Acting Deputy Commissioner, U.S. Customs and Border Protection.

Approved:

Timothy E. Skud,

Deputy Assistant Secretary of the Treasury. [FR Doc. 2018–17847 Filed 8–17–18; 8:45 am]

BILLING CODE 9111-14-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1

[Docket No. FDA-2012-D-1002]

Supplemental Questions and Answers Regarding Food Facility Registration; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the availability of a draft guidance for industry entitled 'Supplemental Questions and Answers Regarding Food Facility Registration." This draft guidance is intended to supplement the guidance document entitled "Questions and Answers Regarding Food Facility Registration." DATES: Submit either electronic or written comments on the draft guidance by October 19, 2018 to ensure that the Agency considers your comment on the draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance 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 that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on https://www.regulations.gov.
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- Mail/Hand delivery/Courier (for written/paper submissions): Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA—2012—D—1002 for "Supplemental Questions and Answers Regarding Food Facility Registration." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at https://www.regulations.gov or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.