■ 7. Supplement no. 4 to part 744 is amended by revising the introductory text to read as follows:

Supplement No. 4 to Part 744—Entity List

This supplement lists certain entities subject to license requirements for specified items under this part 744 and part 746 of the EAR. License requirements for these entities include exports, reexports, and transfers (in-country) unless otherwise stated. A license is required, to the extent specified on the Entity List, to export, reexport, or transfer (in-country) any item subject to the EAR when an entity that is listed on the Entity List is a party to the transaction as described in § 748.5(c) through (f). of the EAR This list of entities is revised and updated on a periodic basis in this Supplement by adding new or amended notifications and deleting notifications no longer in effect.

* * * * *

PART 772—DEFINITIONS OF TERMS

■ 8. The authority citation for 15 CFR part 772 continues to read as follows:

Authority: 50 U.S.C. 4801–4852; 50 U.S.C. 4601 *et seq.;* 50 U.S.C. 1701 *et seq.;* E.O. 13222, 66 FR 44025, 3 CFR, 2001 Comp., p. 783.

■ 9. Section 772.1 is amended by
■ a. Revising the definition of ''Standards-related activity.''

The revisions read as follows:

§ 772.1 Definitions of Terms as Used In the Export Administration Regulations (EAR).

* * * * *

Standards-related activity. See § 734.10 of the EAR.

* * * * *

Thea D. Rozman Kendler,

Assistant Secretary for Export Administration.

[FR Doc. 2024–15810 Filed 7–17–24; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 630

[Docket No. FDA–2022–D–0362]

Blood Pressure and Pulse Donor Eligibility Requirements: Compliance Policy; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance entitled ''Blood Pressure and Pulse Donor Eligibility Requirements: Compliance Policy; Guidance for Industry.'' The guidance document addresses certain regulatory requirements for determining donor eligibility that apply to blood establishments that collect blood and blood components for transfusion or for further manufacturing use, including Source Plasma. In a final rule dated May 22, 2015, FDA amended the regulations applicable to blood establishments for determining donor eligibility and testing blood and blood components. The revised requirements were implemented in order to assure the safety of the blood supply and to protect donor health. This guidance finalizes the draft guidance entitled ''Blood Pressure and Pulse Donor Eligibility Requirements: Compliance Policy; Draft Guidance for Industry'' issued on May 24, 2022.

DATES: The announcement of the guidance is published in the **Federal Register** on July 18, 2024.

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 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–2022–D–0362 for ''Blood Pressure and Pulse Donor Eligibility Requirements: Compliance Policy; Guidance for Industry.'' 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, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management

Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the guidance to the Office of Communication, Outreach and Development, Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 3128, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist the office in processing your requests. The guidance may also be obtained by mail by calling CBER at 1–800–835–4709 or 240–402–8010. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

FOR FURTHER INFORMATION CONTACT: Myrna Hanna, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 7301, Silver Spring, MD 20993–0002, 240–402–7911.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a document entitled ''Blood Pressure and Pulse Donor Eligibility Requirements: Compliance Policy.'' The document addresses certain regulatory requirements for determining donor eligibility that apply to blood establishments that collect blood components for transfusion or for further manufacturing use, including Source Plasma. In the final rule dated May 22, 2015 (80 FR 29841) entitled ''Requirements for Blood and Blood Components Intended for Transfusion or for Further Manufacturing Use,'' FDA amended the regulations applicable to blood establishments for determining donor eligibility and testing blood and blood components.[1] The revised requirements were implemented in order to assure the safety of the blood supply and to protect donor health. The final rule became effective on May 23, 2016. FDA has developed the document in response to feedback from blood establishments regarding the donor eligibility requirements for blood pressure and pulse in 21 CFR 630.10 and the corresponding requirements for medical supervision in 21 CFR 630.5. The guidance describes the circumstances in which FDA does not intend to take regulatory action for a blood establishment's failure to comply with certain regulations for determining the eligibility of blood donors with blood pressure or pulse measurements outside of the specified limits.

This guidance finalizes the draft guidance entitled ''Blood Pressure and Pulse Donor Eligibility Requirements: Compliance Policy; Draft Guidance for Industry'' issued on May 24, 2022 (87 FR 31567). Changes made from the draft to the final guidance took into consideration comments received. After considering the comments, we made a few clarifying edits to the guidance and other editorial changes.

This guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The guidance represents the current thinking of FDA on blood pressure and pulse donor eligibility requirements and explains our compliance policy with respect to these requirements. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

II. Paperwork Reduction Act of 1995

While this guidance contains no collection of information, it does refer to previously approved FDA collections of information. The previously approved collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3521). The collections of information in 21 CFR part 601 have been approved under OMB control number 0910–0338; the collections of information in 21 CFR parts 606 and 630 have been approved under OMB control number 0910–0116.

III. Electronic Access

Persons with access to the internet may obtain the guidance at *https://www.fda.gov/vaccines-blood-biologics/guidance-compliance-regulatory-information-biologics/biologics-guidances, https://www.fda.gov/regulatory-information/search-fda-guidance-documents,* or *https://www.regulations.gov.*

Dated: July 5, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–15228 Filed 7–17–24; 8:45 am]

BILLING CODE 4164–01–P

[1] The Office of the Federal Register has published this document under the category ''Rules and Regulations'' pursuant to 1 CFR 5.9(b). The categorization is solely for purposes of publication in the **Federal Register** and does not change the nature of the document and is not intended to affect its validity, content, or intent. See 1 CFR 5.1(c).

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Part 1

[TD 10004]

RIN 1545–BM19

Guidance Under Section 367(b) Related to Certain Triangular Reorganizations and Inbound Nonrecognition Transactions

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Final regulations.

SUMMARY: This document contains final regulations regarding the treatment of property used to acquire parent stock or securities in connection with certain triangular reorganizations involving one or more foreign corporations; the consequences to persons that receive parent stock or securities pursuant to such reorganizations; and the treatment of certain subsequent inbound nonrecognition transactions following such reorganizations and certain other transactions. The final regulations affect corporations engaged in certain triangular reorganizations involving one or more foreign corporations, certain shareholders of foreign corporations acquired in such reorganizations, and foreign corporations that participate in certain inbound nonrecognition transactions.

DATES:

Effective date: These regulations are effective on July 17, 2024.

Applicability dates: For dates of applicability, *see* §§ 1.367(a)–3(g)(1)(viii), 1.367(b)–3(g)(7)(i), 1.367(b)–4(i), 1.367(b)–6(a)(1)(v) and (vi), 1.367(b)–10(e)(2), (3), and (5), and 1.1411–10(i).

FOR FURTHER INFORMATION CONTACT: Brady Plastaras at (202) 317–6937 (not a toll-free number).

SUPPLEMENTARY INFORMATION:

Background

On October 6, 2023, the Department of the Treasury (Treasury Department) and the IRS published proposed regulations (REG–117614–14) in the **Federal Register** (88 FR 69559) under section 367(b) of the Internal Revenue Code (the ''Proposed Regulations'') that would implement the regulations announced and described in Notice 2014–32 (2014–20 IRB 1006) and Notice 2016–73 (2016–52 IRB 908), with modifications. This document finalizes the Proposed Regulations without substantive change. Terms used but not defined in this preamble have the