Commission informed about the status of the transfer of the rights and assets to Elanco and Bayer.

The Commission's goal in evaluating possible purchasers of divested rights and assets is to maintain the competitive environment that existed prior to the Proposed Acquisition. If the Commission determines that either buyer is not an acceptable acquirer, or that the manner of the divestiture is not acceptable, the proposed Order requires the parties to unwind the sale and then divest the products to another Commission-approved acquirer within six months of the date that the proposed Order becomes final. The proposed Order further allows the Commission to appoint a trustee in the event the parties fail to divest the products.

The purpose of this analysis is to facilitate public comment on the proposed Consent Agreement, and it is not intended to constitute an official interpretation of the proposed Order or to modify its terms in any way.

By direction of the Commission.

April J. Tabor,

Acting Secretary.

[FR Doc. 2016–31848 Filed 1–3–17; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice Designating State Title IV–D Child Support Agencies as "Public Bodies"

AGENCY: Office of Child Support Enforcement, Administration for Children and Families, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: This notice designates state IV–D child support agencies as public bodies authorized to perform specific functions of the Central Authority under Article 6(3) of the the Hague Convention of 23 November 2007 on the International Recovery of Child Support and Other Forms of Family Maintenance (Convention).and specifies functions to be performed by the state agencies in relation to applications under the Convention.

ADDRESSES: Interested parties may submit written comments on this notice to the United States Central Authority for International Child Support, Department of Health and Human Services, Office of Child Support Enforcement, 330 C Street SW., 5th Floor, Washington, DC 20201. Comments received will be available for public inspection at this address from 9:00 a.m. to 5:00 p.m. EST, Monday through Friday.

DATES: The Convention will enter into force for the United States on January 1, 2017.

FOR FURTHER INFORMATION CONTACT: The Division of Policy and Training, Office of Child Support Enforcement, Administration for Children and Families, 330 C Street SW., 5th Floor, Washington, DC 20201.

SUPPLEMENTARY INFORMATION: The President signed the Instrument of Ratification on August 30, 2016, and the United States of America deposited its Instrument of Ratification of the Convention on September 7, 2016. The Convention will enter into force for the United States on January 1, 2017. Section 459A of the Social Security Act (42 U.S.C. 659a) and Executive Order 13752, 81 FR 90181 (Dec. 8, 2016) designate the Department of Health and Human Services as the Central Authority of the United States for purposes of the Convention, and authorize the Secretary of Health and Human Services to perform all lawful acts that may be necessary and proper in order to execute the functions of the Central Authority. Article 6(3) of the Convention authorizes the designation of public bodies to perform specific functions under the Convention, subject to the supervision of the Central Authority. The Executive Order specifically authorizes the designation of the state agencies responsible for implementing an approved State Plan under title IV–D of the Social Security Act, 42 U.S.C. 651 *et seq.,* as public bodies authorized to perform specific functions in relation to applications under the Convention. All states have enacted the Uniform Interstate Family Support Act of 2008 (UIFSA 2008) to enable uniform implementation of the Convention in the United States.

Under authority delegated by the Secretary for administration of the title IV–D program, I hereby designate the state title IV–D child support agencies as public bodies authorized to perform functions related to applications under the Convention in accordance with UIFSA 2008, title IV–D and title IV–D regulations, and guidance and instructions, subject to the supervision of the federal Office of Child Support Enforcement. Such functions shall include the provision of support enforcement services to applicants under the Convention, including: Transmitting and receiving applications under the Convention; initiating or facilitating the institution of proceedings with respect to applications; establishing paternity and support orders; recognizing, modifying, and enforcing such orders; collecting and distributing payments under such orders; and providing administrative and legal services without cost to applicants.

Statutory Authority: Section 459(a) of the Social Security Act (42 U.S.C. 659(a)

Dated: December 29, 2016.

Mark H. Greenberg,

Acting Assistant Secretary for Children and Families.

[FR Doc. 2016–31895 Filed 1–3–17; 8:45 am]

BILLING CODE 4184–42–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–0001]

Advisory Committee; Technical Electronic Product Radiation Safety Standards Committee, Renewal

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; renewal of advisory committee.

SUMMARY: The Food and Drug Administration (FDA) is announcing the renewal of the Technical Electronic Product Radiation Safety Standards Committee by the Commissioner of Food and Drugs (the Commissioner). The Commissioner has determined that it is in the public interest to renew the Technical Electronic Product Radiation Safety Standards Committee for an additional 2 years beyond the charter expiration date. The new charter will be in effect until December 24, 2018.

DATES: Authority for the Technical Electronic Product Radiation Safety Standards Committee will expire on December 24, 2016, unless the Commissioner formally determines that renewal is in the public interest.

FOR FURTHER INFORMATION CONTACT: Shanika Craig, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1613, Silver Spring, MD, 20993–0002, 301–796–6639, *Shanika.Craig@fda.hhs.gov*

SUPPLEMENTARY INFORMATION: Pursuant to 41 CFR 102–3.65 and approval by the Department of Health and Human Services pursuant to 45 CFR part 11 and by the General Services Administration, FDA is announcing the renewal of the Technical Electronic Product Radiation Safety Standards Committee. The

committee is a non-discretionary Federal advisory committee established to provide advice and consultation to the Commissioner. The Commissioner of Food and Drugs is charged with the administration of the Radiation Control for Health and Safety Act of 1968. This Act creates the Technical Electronic Product Radiation Safety Standards Committee and requires the Commissioner to consult with the Committee before prescribing standards for radiation emissions from electronic products. This Committee provides advice and consultation to the Commissioner of Food and Drugs on the technical feasibility, reasonableness, and practicability of performance standards for electronic products to control the emission of radiation from such products, and may recommend electronic product radiation safety standards to the Commissioner for consideration.

The Committee shall consist of a core of 15 voting members including the Chair. Members and the Chair are selected by the Commissioner or designee from among authorities knowledgeable in the fields of science or engineering applicable to electronic product radiation safety. Members will be invited to serve for overlapping terms of up to 4 years. Terms of more than two years are contingent upon the renewal of the Committee by appropriate action prior to its expiration. The core of voting members will include five members selected from governmental agencies, including State and Federal Governments, five members from the affected industries, and five members from the general public, of which at least one shall be a representative of organized labor. A quorum shall consist of 10 members, of which at least 3 shall be from the general public, 3 from the government agencies, and 3 from the affected industries.

Further information regarding the most recent charter and other information can be found at *http://www.fda.gov/AdvisoryCommittees/CommitteesMeetingMaterials/Radiation-EmittingProducts/TechnicalElectronicProductRadiationSafetyStandardsCommittee/default.htm.* or by contacting the Designated Federal Officer (see **FOR FURTHER INFORMATION CONTACT**). In light of the fact that no change has been made to the committee name or description of duties, no amendment will be made to 21 CFR 14.100.

This document is issued under the Federal Advisory Committee Act (5 U.S.C. app.). For general information related to FDA advisory committees, please visit us at *http://www.fda.gov/AdvisoryCommittees/default.htm.*

Dated: December 28, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–31847 Filed 1–3–17; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–D–4437]

In-Use Stability Studies and Associated Labeling Statements for Multiple-Dose Injectable Animal Drug Products; Draft 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 draft guidance for industry (GIF) #242 entitled ''In-Use Stability Studies and Associated Labeling Statements for Multiple-Dose Injectable Animal Drug Products.'' The purpose of in-use stability testing is to establish a period of time during which a multiple-dose drug product may be used while retaining acceptable quality specifications once the container is opened (*e.g.,* after a container has been needle-punctured). This draft guidance reflects the Agency's current thinking on how to formulate in-use statements, as well as how to design and carry out in-use stability studies to support these in-use statements, for multiple-dose injectable drug products intended for use in animals. This current thinking pertains to both generic drug products and pioneer drug products regardless of whether or not the pioneer reference listed new animal drug (RLNAD) currently has an in-use statement on the labeling.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by March 6, 2017.

ADDRESSES: You may submit comments 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):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, 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–2016–D–4437 for ''In-Use Stability Studies and Associated Labeling Statements for Multiple-Dose Injectable Animal Drug Products.'' 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 Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- 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