

DEPARTMENT OF AGRICULTURE**Animal and Plant Health Inspection Service****9 CFR Part 112**

[Docket No. APHIS–2011–0049]

RIN 0579–AD64

Viruses, Serums, Toxins, and Analogous Products; Single Label Claim for Veterinary Biological Products**AGENCY:** Animal and Plant Health Inspection Service, USDA.**ACTION:** Final rule; technical amendment.

SUMMARY: In a final rule published in the *Federal Register* on July 10, 2015, and effective on September 8, 2015, we amended the Virus-Serum-Toxin Act regulations to provide for the use of a simpler labeling format that would better communicate product performance to the user. Among other things, we provided the address of a Web site for accessing transmittal forms to be used with each submission of sketches and labels. However, the Web site address provided is incorrect. Therefore, we are amending the regulations to provide the correct address.

DATES: Effective July 20, 2016.**FOR FURTHER INFORMATION CONTACT:** Dr. Donna Malloy, Operational Support Section, Center for Veterinary Biologics, Policy, Evaluation, and Licensing, VS, APHIS, 4700 River Road, Unit 148, Riverdale, MD 20737; (301) 851–2352.

SUPPLEMENTARY INFORMATION: In a final rule¹ that was published in the *Federal Register* on July 10, 2015 (80 FR 39669–39675, Docket No. APHIS–2011–0049), and effective on September 8, 2015, we amended the Virus-Serum-Toxin Act regulations to provide for the use of a simpler labeling format that would better communicate product performance to the user. Among other things, we provided the address of a Web site in § 112.5(a) for accessing transmittal forms to be used with each submission of sketches (including proofs) and labels. However, the Web site address provided is for accessing product licensing data and not transmittal forms. Therefore, we are amending § 112.5(a) to correct the address.

¹ To view the final rule and supporting documents, go to <http://www.regulations.gov/#/docketDetail;D=APHIS-2011-0049>.

Lists of Subjects in 9 CFR Part 112

Animal biologics, Exports, Imports, Labeling, packaging and containers, Reporting and recordkeeping requirements.

Accordingly, we are amending 9 CFR part 112 as follows:

PART 112—PACKAGING AND LABELING

■ 1. The authority citation for part 112 continues to read as follows:

Authority: 21 U.S.C. 151–159; 7 CFR 2.22, 2.80, and 371.4.

§ 112.5 [Amended]

■ 2. In § 112.5, paragraph (a) is amended by removing the words “(productdata.aphis.usda.gov)” and adding the words “(http://www.aphis.usda.gov/animalhealth/cvb/forms)” in their place.

Done in Washington, DC, this 14th day of July 2016.

Jere L. Dick,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2016–17073 Filed 7–19–16; 8:45 am]

BILLING CODE 3410–34–P**NUCLEAR REGULATORY COMMISSION****10 CFR Part 2**

[NRC–2016–0117]

RIN 3150–AJ76

Update to Transcript Correction Procedures**AGENCY:** Nuclear Regulatory Commission.**ACTION:** Final rule.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) is amending its regulation that governs the correction of official transcripts for agency adjudicatory proceedings. The current regulation has not been substantively updated since it was adopted in 1962 and the NRC’s internal procedures have evolved since that time to incorporate technological development. The NRC is not soliciting public comment on this change because the change is limited to an agency rule of procedure and practice that does not affect the rights and responsibilities of outside parties.

DATES: This final rule is effective on July 20, 2016.

ADDRESSES: Please refer to Docket ID NRC–2016–0117 when contacting the NRC about the availability of information for this action. You may

obtain publicly-available information related to this action by any of the following methods:

- Federal Rulemaking Web site: Go to <http://www.regulations.gov> and search for Docket ID NRC–2016–0117. Address questions about NRC dockets to Carol Gallagher; telephone: 301–415–3463; email: Carol.Gallagher@nrc.gov. For other questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.

- NRC’s Public Document Room (PDR): You may examine and purchase copies of public documents at the NRC’s PDR, Room O1–F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852.

FOR FURTHER INFORMATION CONTACT: Tison Campbell, Office of the General Counsel, telephone: 301–287–9290, email: Tison.Campbell@nrc.gov, U.S. Nuclear Regulatory Commission, Washington DC 20555–0001.

SUPPLEMENTARY INFORMATION:**I. Summary of Changes**

In 1962, the Atomic Energy Commission (the NRC’s predecessor agency) adopted revised rules of practice and procedure to govern the conduct of adjudicatory proceedings before the agency (27 FR 377; January 13, 1962). As part of those regulations, the Commission adopted a paragraph governing the correction of hearing transcripts. That provision, originally at § 2.750(b) of title 10 of the *Code of Federal Regulations* (10 CFR), provided specific, prescriptive direction to the Commission’s staff regarding the method for recording and showing corrections to transcripts. For example, the Secretary was directed to make any physical corrections to the official transcript, not by replacing pages, but by drawing a line through the text to be changed in the original transcript and writing the correct text immediately above.

The current agency practice varies. In Commission proceedings, an appendix listing the transcript corrections and a clean version of the transcript are attached to the order adopting the parties’ proposed transcript corrections. In Atomic Safety and Licensing Board Panel proceedings, the boards generally issue an order adopting the parties’ joint proposed transcript corrections, with or without an appendix listing the corrections. The Secretary does not prepare transcripts of board proceedings.

The NRC is, therefore, updating the regulation that governs the correction of official transcripts for agency adjudicatory proceedings, currently at