DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2013–F–1539]

DSM Nutritional Products, Inc.; Withdrawal of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification; withdrawal of petition for rulemaking.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the withdrawal, without prejudice to a future filing, of a food additive petition (animal use) (FAP 2276) proposing that the food additive regulations be amended to provide for the safe use of ethoxyquin in vitamin D formulations, including 25-hydroxyvitamin D_3, used in animal food.

DATES: The food additive petition was withdrawn on September 13, 2017.

ADDRESSES: For access to the docket to read background documents or 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.

FOR FURTHER INFORMATION CONTACT: Chelsea Trull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–6729, *chelsea.trull@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of December 23, 2013 (78 FR 77384), we announced that we had filed a food additive petition (FAP 2276), submitted by DSM Nutritional Products, 45 Waterview Blvd., Parsippany, NJ 07054. The petition proposed to amend the food additive regulations in 21 CFR part 573 *Food Additives Permitted in Feed and Drinking Water of Animals* to provide for the safe use of ethoxyquin as a chemical preservative in vitamin D formulations, including 25-hydroxyvitamin D_3, used in animal food. DSM Nutritional Products, Inc., has now withdrawn the petition without prejudice to a future filing (21 CFR 571.7).

Dated: October 26, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–23728 Filed 10–31–17; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2014–F–0295]

DSM Nutritional Products, Inc.; Withdrawal of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification; withdrawal of petition for rulemaking.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the withdrawal, without prejudice to a future filing, of a food additive petition (FAP 2280) proposing that the food additive regulations be amended to provide for the safe use of 25-hydroxyvitamin D_3 in feed for swine.

DATES: The food additive petition was withdrawn on September 13, 2017.

ADDRESSES: For access to the docket to read background documents or 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.

FOR FURTHER INFORMATION CONTACT: Chelsea Trull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–6729, *chelsea.trull@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of March 25, 2014 (79 FR 16252), we announced that we had filed a food additive petition (FAP 2280), submitted by DSM Nutritional Products, 45 Waterview Blvd., Parsippany, NJ 07054. The petition proposed to amend the food additive regulations in 21 CFR part 573, *Food Additives Permitted in Feed and Drinking Water of Animals,* to provide for the safe use of 25-hydroxyvitamin D_3 in feed for swine. DSM Nutritional Products, Inc. has now withdrawn the petition without prejudice to a future filing (21 CFR 571.7).

Dated: October 26, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–23729 Filed 10–31–17; 8:45 am]

BILLING CODE 4164–01–P

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Part 1

[MD Docket Nos. 17–134; FCC 17–111]

Assessment and Collection of Regulatory Fees for Fiscal Year 2017

AGENCY: Federal Communications Commission.

ACTION: Notice of proposed rulemaking.

SUMMARY: In this document, the Federal Communications Commission (Commission) seeks further comment on the appropriate tiers for calculating terrestrial and satellite international bearer circuit fees, and the methodology by which cable television subscribers in multiple dwelling units (MDUs) are calculated.

DATES: Comments are due on or before December 1, 2017 and reply comments are due on or before December 18, 2017.

ADDRESSES: You may submit comments, identified by MD Docket No. 17–134, by any of the following methods listed in the Comment Filing Procedures section below.

For detailed instructions for submitting comments and additional information on the rulemaking process, see the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: Roland Helvajian, Office of Managing Director at (202) 418–0444.

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's *Further Notice of Proposed Rulemaking* (FNPRM), FCC 17–111, MD Docket No. 17–134 adopted on September 1, 2017 and released on September 5, 2017. The full text of this document is available for inspection and copying during normal business hours in the FCC Reference Center, 445 12th Street SW., Room CY–A257, Portals II, Washington, DC 20554, and may also be purchased from the Commission's copy contractor, BCPI, Inc., Portals II, 445 12th Street SW., Room CY–B402, Washington, DC 20554. Customers may contact BCPI, Inc. via their Web site, *http://www.bcpi.com,* or call 1–800–378–3160. This document is available in alternative formats (computer diskette, large print, audio record, and Braille). Persons with disabilities who need documents in