Commission's swap data reporting rules. I commend the cross-divisional data team's effort to fix our reporting rules and enhance the Commission's ability to use its data. I hope that the data team and the Commission will carefully evaluate market participants' comments and recommendations and develop workable solutions to improve our data reporting regime.

At the same time, I urge market participants to carefully review the Commission's questions, submit their comments, and alert the Commission to other data reporting issues that have not been included in this request for comment. This comment period is a critical step in the Commission's effort to improve its data utilization. I encourage all market participants to help the Commission improve its data reporting regime.

[FR Doc. 2014–06426 Filed 3–25–14; 8:45 am]

BILLING CODE 6351–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2014–F–0296]

DSM Nutritional Products; Filing of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of petition.

SUMMARY: The Food and Drug Administration (FDA) is announcing that DSM Nutritional Products has filed a petition proposing that the food additive regulations be amended to provide for the safe use of 25-hydroxyvitamin D_3 in feed for turkeys.

DATES: Submit either electronic or written comments on the petitioner's request for categorical exclusion from preparing an environmental assessment or environmental impact statement by April 25, 2014.

ADDRESSES: Submit electronic comments to: *http://www.regulations.gov*. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Isabel W. Pocurull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240-453-6853.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (section 409(b)(5) (21 U.S.C. 348(b)(5)), notice is given that a food additive petition (FAP 2279) has been filed by DSM Nutritional Products, 45 Waterview Blvd., Parsippany, NJ 07054. The petition proposes to amend Title 21 of the Code of Federal Regulations (CFR) in part 573 *Food Additives Permitted in Feed and Drinking Water of Animals* (21 CFR part 573) to provide for the safe use of 25-hydroxyvitamin D_3 in feed for turkeys.

The petitioner has requested a categorical exclusion from preparing an environmental assessment or environmental impact statement under 21 CFR 25.32(r). Interested persons may submit either electronic or a single copy of written comments regarding this request for categorical exclusion to the Division of Dockets Management (see **DATES** and **ADDRESSES**). Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: March 21, 2014.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

[FR Doc. 2014–06623 Filed 3–25–14; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Chapter I

[Docket No. FDA–2013–N–0590]

RIN 0910–AG97

Implementation of the Food and Drug Administration Food Safety Modernization Act Amendments to the Reportable Food Registry Provisions of the Federal Food, Drug, and Cosmetic Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Advance notice of proposed rulemaking.

SUMMARY: The Food and Drug Administration (FDA) is issuing this advance notice of proposed rulemaking (ANPRM) to solicit comments, data, and information to assist the Agency in implementing the FDA Food Safety Modernization Act (FSMA), which added new provisions to the Reportable Food Registry (RFR) requirements of the Federal Food, Drug, and Cosmetic Act (the FD&C Act). Under the new provisions, FDA may require a responsible party to also submit to FDA ''consumer-oriented'' information regarding certain reportable foods, including information necessary to enable a consumer to accurately identify whether the consumer is in possession of a reportable food. FDA must prepare and publish on FDA's Internet Web site a one-page summary of the consumer-oriented information that can be easily printed by a grocery store for the purposes of consumer notification. A grocery store that sold a reportable food that is the subject of an FDA one-page summary, and that is part of a chain of establishments with 15 or more physical locations, is required to prominently display the FDA one-page summary, or the information from the summary, within 24 hours after the one-page summary is published on FDA's Web site, through a method identified by FDA. FDA is seeking input on topics including consumer-oriented information submissions, consumer notifications, posting consumer notifications in grocery stores, and grocery stores subject to the new requirements.

DATES: Submit either electronic or written comments by June 9, 2014.

ADDRESSES: You may submit comments, identified by Docket No. FDA–2013–N–0590 or Regulatory Information Number (RIN) number 0910–AG97, by any of the following methods:

Electronic Submissions

Submit electronic comments in the following way:

- Federal eRulemaking Portal: *http://www.regulations.gov*. Follow the instructions for submitting comments.

Written Submissions

Submit written submissions in the following ways:

- Mail/Hand delivery/Courier (for paper submissions): Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Instructions: All submissions received must include the Agency name and Docket No. FDA–2013–N–0590 and RIN 0910–AG97 for this advance notice of proposed rulemaking. All comments received may be posted without change to *http://www.regulations.gov,* including any personal information provided. For additional information on submitting comments, see the ''Comments'' heading of the **SUPPLEMENTARY INFORMATION** section of this document.

Docket: For access to the docket to read background documents or comments received, go to *http://www.regulations.gov* and insert the docket number(s), found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Division of Dockets