

25 August 2025

Original: English

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## **Committee on Sanitary and Phytosanitary Measures**

## **NOTIFICATION**

1.	Notifying Member: <u>UNITED STATES OF AMERICA</u> If applicable, name of local government involved:
2.	Agency responsible: Food and Drug Administration (FDA)
3.	Products covered (provide tariff item number(s) as specified in national schedules deposited with the WTO; ICS numbers should be provided in addition, where applicable): Preparations of a kind used in animal feeding (HS code(s): 2309); Food technology (ICS code(s): 67)
4.	Regions or countries likely to be affected, to the extent relevant or practicable:
	[X] All trading partners
	[ ] Specific regions or countries:
5.	<b>Title of the notified document:</b> Evonik Corporation; Filing of Food Additive Petition (Animal Use); Notification of Petition. <b>Language(s):</b> English. <b>Number of pages:</b> 1
	https://www.federalregister.gov/d/2025-15792 https://members.wto.org/crnattachments/2025/SPS/USA/25_05523_00_e.pdf
6.	<b>Description of content:</b> The Food and Drug Administration (FDA or we) is announcing that we have filed a food additive petition, submitted by Evonik Corporation, proposing that we amend our food additive regulations to provide for the safe use of <i>Bacillus velezensis</i> as a source of viable microorganism in animal food for all species.
	The food additive petition was filed on 18 July 2025.
7.	Objective and rationale: [X] food safety, [X] animal health, [ ] plant protection, [ ] protect humans from animal/plant pest or disease, [ ] protect territory from other damage from pests.
8.	Is there a relevant international standard? If so, identify the standard:
	[ ] Codex Alimentarius Commission (e.g. title or serial number of Codex standard or related text):
	[] World Organization for Animal Health (OIE) (e.g. Terrestrial or Aquatic Animal Health Code, chapter number):
	[ ] International Plant Protection Convention (e.g. ISPM number):
	[X] None
	Does this proposed regulation conform to the relevant international standard?
	[ ] Yes [ ] No
	If no, describe, whenever possible, how and why it deviates from the international standard:

- 9. Other relevant documents and language(s) in which these are available:
- 10. Proposed date of adoption (dd/mm/yy): Not applicable
  Proposed date of publication (dd/mm/yy): Not applicable
- 11. Proposed date of entry into force: [ ] Six months from date of publication, and/or (dd/mm/yy): Not applicable
  - [X] Trade facilitating measure
- 12. Final date for comments: [ ] Sixty days from the date of circulation of the notification and/or (dd/mm/yy): Not applicable

Agency or authority designated to handle comments: [ ] National Notification Authority, [ ] National Enquiry Point. Address, fax number and e-mail address (if available) of other body:

For access to the docket to read background documents or comments received, go to <a href="https://www.regulations.gov">https://www.regulations.gov</a> 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: Megan Hall, Center for Veterinary Medicine, Food and Drug Administration, 5001 Campus Drive, College Park, MD 20740, +(301) 796 3801, Megan.Hall@fda.hhs.gov

13. Text(s) available from: [ ] National Notification Authority, [ ] National Enquiry Point. Address, fax number and e-mail address (if available) of other body:

Text can be found in the Federal Register, Vol. 90, No. 158, Page 40269 or on the Internet at: <a href="https://www.federalregister.gov/d/2025-15792">https://www.federalregister.gov/d/2025-15792</a>