

13 September 2022

Original: English

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Committee on Sanitary and Phytosanitary Measures

NOTIFICATION

- 1. Notifying Member: <u>EUROPEAN UNION</u>

 If applicable, name of local government involved:
- 2. Agency responsible: European Commission, Health and Food Safety Directorate-General
- 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)
- 4. Regions or countries likely to be affected, to the extent relevant or practicable:
 - [X] All trading partners
 - [] Specific regions or countries:
- 5. Title of the notified document: Commission Implementing Regulation (EU) 2022/1493 of 8 September 2022 concerning the authorisation of L-methionine produced by Corynebacterium glutamicum KCCM 80245 and Escherichia coli KCCM 80246 as feed additives for all animal species (Text with EEA relevance). Language(s): English, French and Spanish. Number of pages: 5

https://members.wto.org/crnattachments/2022/SPS/EEC/22 6087 00 e.pdf https://members.wto.org/crnattachments/2022/SPS/EEC/22 6087 00 f.pdf https://members.wto.org/crnattachments/2022/SPS/EEC/22 6087 00 s.pdf

- 6. Description of content: In accordance with Article 7 of Regulation (EC) No 1831/2003 an application was submitted for the authorisation of L-methionine produced by Corynebacterium glutamicum KCCM 80245 and Escherichia coli KCCM 80246. That application was accompanied by the particulars and documents required under Article 7 of that Regulation. The application concerns the authorisation of L-methionine produced by Corynebacterium glutamicum KCCM 80245 and Escherichia coli KCCM 80246 as feed additives for all animal species to be classified in the additive category 'nutritional The Authority further concluded that L-methionine produced Corynebacterium glutamicum KCCM 80245 and Escherichia coli KCCM 80246 is an effective source of methionine for all animal species and that in order to be as efficacious in ruminants as in non-ruminant species, the substance should be protected against degradation in the rumen. The Authority does not consider that there is a need for specific requirements of post-market monitoring. It also verified the report on the method of analysis of the feed additives in feed submitted by the Reference Laboratory set up by Regulation (EC) No 1831/2003.
- 7. Objective and rationale: [X] food safety, [] 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): Code of practice on Good Animal Feeding CAC/RCP 54-2004 World Organization for Animal Health (OIE) (e.g. Terrestrial or Aquatic [] Animal Health Code, chapter number): International Plant Protection Convention (e.g. ISPM number): [] [] Does this proposed regulation conform to the relevant international standard? [X] 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): 8 September 2022 Proposed date of publication (dd/mm/yy): 9 September 2022 11. Proposed date of entry into force: [] Six months from date of publication, and/or (dd/mm/yy): This Regulation shall enter into force on the twentieth day following its publication in the Official Journal of the European Union. [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: [X] National Notification Authority, [X] National Enquiry Point. Address, fax number and e-mail address (if available) of other body:

European Commission

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

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