NOTIFICATION

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| **1.** | **Notifying Member:** European Union**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):** Preparation of a kind used in animal nutrition (HS Code: 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) 2021/2076 of 26 November 2021 concerning the authorisation of L-tryptophan produced by *Escherichia coli* KCCM 80210 as a feed additive for all animal species (Text with EEA relevance).**Language(s):** English, French and Spanish. **Number of pages:** 4<https://members.wto.org/crnattachments/2021/SPS/EEC/21_7476_00_e.pdf><https://members.wto.org/crnattachments/2021/SPS/EEC/21_7476_00_f.pdf><https://members.wto.org/crnattachments/2021/SPS/EEC/21_7476_00_s.pdf> |
| **6.** | **Description of content:** Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation. In accordance with Article 7 of Regulation (EC) No 1831/2003 an application was submitted for the authorisation of L-tryptophan produced by *Escherichia coli* KCCM 80210. The application was accompanied by the particulars and documents required under Article 7 of that Regulation. The application concerns the authorisation of L‑tryptophan produced by *Escherichia coli* KCCM 80210 as a feed additive for all animal species, to be classified in the additive category 'nutritional additives', functional group 'amino acids, their salts and analogues'. The European Food Safety Authority ('the Authority') concluded in its opinion of 27 January 2021 that, under the proposed conditions of use, L-tryptophan produced by *Escherichia coli* KCCM 80210 does not have an adverse effect on the health of non-ruminant animals, consumer safety or the environment. To be safe for ruminants, the L-tryptophan should be protected against degradation in the rumen. The Authority stated that the additive under assessment is considered a mild eye irritant. The endotoxin activity of the additive and its dusting potential indicate a risk by inhalation. Therefore, the Commission considers that appropriate protective measures should be taken to prevent adverse effects on human health, in particular as regards the users of the additive. |
| **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:****[****X]** **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)*:** **[ ]****None****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)*:** 26 November 2021**Proposed date of publication *(dd/mm/yy)*:** 29 November 2021 |
| **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 CommissionDG Health and Food Safety, Unit D2-Multilateral International RelationsRue Froissart 101B-1049 BrusselsTel: +(32 2) 29 54263Fax: +(32 2) 29 98090E-mail: sps@ec.europa.eu |
| **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:** European CommissionDG Health and Food Safety, Unit D2-Multilateral International RelationsRue Froissart 101B-1049 BrusselsTel: +(32 2) 29 54263Fax: +(32 2) 29 98090E-mail: sps@ec.europa.eu |