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) 2021/2092 concerning the authorisation of potassium diformate as a feed additive for pigs for fattening and weaned piglets (Text with EEA relevance).**Language(s):** English, French and Spanish. **Number of pages:** 3<https://members.wto.org/crnattachments/2021/SPS/EEC/21_7414_00_e.pdf><https://members.wto.org/crnattachments/2021/SPS/EEC/21_7414_00_f.pdf><https://members.wto.org/crnattachments/2021/SPS/EEC/21_7414_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 an authorisation. In accordance with Article 7 of Regulation (EC) No 1831/2003 an application was submitted for the authorisation of potassium diformate. That application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003. The application concerns the authorisation of potassium diformate as a feed additive for pigs for fattening and weaned piglets, to be classified in the category 'technological additives'. The European Food Safety Authority ('the Authority') concluded in its opinion of 5 May 2021 that, under the proposed conditions of use, potassium diformate does not have adverse effects on animal health, consumer safety or the environment. It also concluded that the substance does not raise concerns regarding the effects on the respiratory system and the skin but is an eye irritant. 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. The Authority also concluded that the substance has the potential to be efficacious as a technological additive in feedingstuffs. It also verified the report on the methods of analysis of the feed additive 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:****[****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)*:** 29 November 2021**Proposed date of publication *(dd/mm/yy)*:** 30 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 |