

15 July 2020

(20-4819)

Page: 1/3

Committee on Sanitary and Phytosanitary Measures

Original: English

NOTIFICATION

- 1. Notifying Member: <u>EUROPEAN UNION</u>
 - If applicable, name of local government involved:
- 2. Agency responsible: European Commission, Health and Consumers 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): Cereals (HS Codes: 1001, 1002, 1003, 1004, 1005, 1006, 1007, 1008), foodstuffs of animal origin (HS Codes: 0201, 0202, 0203, 0204, 0205, 0206, 0207, 0208, 0209, 0210) and certain products of plant origin, including fruit and vegetables.
- 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: Draft Commission Regulation amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for fluxapyroxad, hymexazol, metamitron, penflufen and spirotetramat in or on certain products (Text with EEA relevance). Language(s): English Number of pages: 33

https://members.wto.org/crnattachments/2020/SPS/EEC/20 4139 01 e.pdf https://members.wto.org/crnattachments/2020/SPS/EEC/20 4139 00 e.pdf https://members.wto.org/crnattachments/2020/SPS/EEC/20 4139 02 e.pdf https://members.wto.org/crnattachments/2020/SPS/EEC/20 4139 03 e.pdf https://members.wto.org/crnattachments/2020/SPS/EEC/20 4139 04 e.pdf

- 6. **Description of content:** The proposed draft Regulation concerns the review of existing MRLs for fluxapyroxad, hymexazol, metamitron, penflufen and spirotetramat in certain food commodities. MRLs for these substances in certain commodities are changed: either increased or lowered. Lower MRLs are set after updating the limits of determination and/or deleting old uses which are not authorised any more in the European Union or for which a human health concern may not be excluded. The residue definition for some substances has also been updated.
- 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.

 standard or relation of the second sec	arius Commission (e.g. title or serial number of Codex ted text): Codex Maximum Residue Limits for fluxapyroxad and Residue Limits for spirotetramat. The list of Codex MRLs is rg/fao-who-codexalimentarius/codex- //pesticides/en/ tion for Animal Health (OIE) (e.g. Terrestrial or Aquatic Code, chapter number): ant Protection Convention (e.g. ISPM number): qulation conform to the relevant international standard? never possible, how and why it deviates from the I: The European Food Safety Authority published reasoned f the existing maximum residue levels for fluxapyroxad and for hese opinions, Regulation (EC) No 396/2005 should be amended ents and language(s) in which these are available: 9.396/2005 of the European Parliament and of the Council of a maximum residue levels of pesticides in or on food and feed of
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	in and amending Council Directive 91/414/EEC" <u>a.eu/legal-content/EN/ALL/?uri=CELEX%3A32005R0396</u> ity Authority; Reasoned opinion on the review of the existing vels for fluxapyroxad according to Article 12 of Regulation (EC) Journal 2020;18(1):5984. <u>opa.eu/fr/efsajournal/pub/5984</u> ity Authority; Reasoned opinion on the review of the existing vels for hymexazol according to Article 12 of Regulation (EC) No rnal 2019;17(11):5895. <u>ropa.eu/en/efsajournal/pub/5895</u> ity Authority; Reasoned opinion on the review of the existing vels for metamitron according to Article 12 of Regulation (EC) Journal 2020;18(1):5959 <u>orary.wiley.com/doi/full/10.2903/j.efsa.2020.5959</u> ity Authority; Reasoned opinion on the review of the existing vels for penflufen according to Article 12 of Regulation (EC) No rnal 2019;17(10):5840. <u>ropa.eu/fr/efsajournal/pub/5840</u> ity Authority; Reasoned opinion on the review of the existing vels for penflufen according to Article 12 of Regulation (EC) No rnal 2019;17(10):5840. <u>ropa.eu/fr/efsajournal/pub/5840</u> ity Authority; Reasoned opinion on the review of the existing vels for spirotetramat according to Article 12 of Regulation (EC) No

10. Proposed date of adoption (*dd/mm/yy*): February 2021.

Proposed date of publication (*dd/mm/yy*): April 2021.

- 3 -

11. Proposed date of entry into force: [] Six months from date of publication, and/or (*dd/mm/yy*): Twenty days after publication in the Official Journal of the European Union.

[] Trade facilitating measure

12. Final date for comments: [X] Sixty days from the date of circulation of the notification and/or (*dd/mm/yy*): 13 September 2020

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 DG Health and Food Safety, Unit D2 - Multilateral international relations Rue Froissart 101 B 1049 Brussels Tel: +(322) 295 4263 Fax: +(322) 299 8090 E-mail: <u>sps@ec.europa.eu</u>

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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