



14 November 2014

(14-6673)

Page: 1/3

Committee on Sanitary and Phytosanitary Measures

Original: English

**NOTIFICATION**

1.	<b>Notifying Member:</b> <u>EUROPEAN UNION</u> <b>If applicable, name of local government involved:</b>
2.	<b>Agency responsible:</b> European Commission, Health and Consumers Directorate-General
3.	<b>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):</b> 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.	<b>Regions or countries likely to be affected, to the extent relevant or practicable:</b> <input checked="" type="checkbox"/> All trading partners <input type="checkbox"/> Specific regions or countries:
5.	<b>Title of the notified document:</b> Annexes to "Draft Regulation amending Annexes II, III and V to Regulation (EC) No. 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 1-methylcyclopropene, flonicamid, flutriafol, indolylacetic acid, indolylbutyric acid, pethoxamid, pirimicarb, prothioconazole and teflubenzuron in or on certain products." <b>Language(s):</b> English <b>Number of pages:</b> 5+79 <a href="http://members.wto.org/crnattachments/2014/sps/EEC/14_5100_00_e.pdf">http://members.wto.org/crnattachments/2014/sps/EEC/14_5100_00_e.pdf</a> <a href="http://members.wto.org/crnattachments/2014/sps/EEC/14_5100_01_e.pdf">http://members.wto.org/crnattachments/2014/sps/EEC/14_5100_01_e.pdf</a>
6.	<b>Description of content:</b> These notified annexes to the draft Regulation set proposed maximum residue levels (MRLs) for 1-methylcyclopropene, flonicamid, flutriafol, indolylacetic acid, indolylbutyric acid, pethoxamid, pirimicarb, prothioconazole and teflubenzuron. MRLs for these substances in certain commodities are changed: either increased or lowered. Lower MRLs are set after updating the limit of determinations and/or deleting old uses which are not authorized any more in the European Union or for which a human health concern may not be excluded.
7.	<b>Objective and rationale:</b> <input checked="" type="checkbox"/> food safety, <input type="checkbox"/> animal health, <input type="checkbox"/> plant protection, <input type="checkbox"/> protect humans from animal/plant pest or disease, <input type="checkbox"/> 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)** Codex Maximum Residue Limits for the following pesticides: 1-methylcyclopropene, flonicamid, flutriafof, indolylacetic acid, indolylbutyric acid, pethoxamid, pirimicarb, prothioconazole and teflubenzuron.

☐ **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?**

☒ 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:**

- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for 1-methylcyclopropene according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(7):3746, 23 pp.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for flonicamid according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(6):3740, 4- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for flutriafof according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(5):3787, 64 pp.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for indolylacetic acid according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(7):3747, 8 pp.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for indolylbutyric acid according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(7):3748, 8 pp.
- Commission Decision 2008/941/EC of 8 December 2008 concerning the non-inclusion of certain active substances in Annex I to Council Directive 91/414/EEC and the withdrawal of authorisations for plant protection products containing these substances. OJ L 335, 13.12.2008, p. 91-93.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for pethoxamid according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(7):3749, 32 pp.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for pirimicarb according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(5):3688, 107 pp.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for prothioconazole according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(5):3689, 72 pp.
- EFSA (European Food Safety Authority), 2014. Reasoned opinion on the review of the existing maximum residue levels (MRLs) for teflubenzuron according to Article 12 of Regulation (EC) No. 396/2005. EFSA Journal 2014; 12(4):3664, 39 pp.

**10. Proposed date of adoption (dd/mm/yy):** March 2015

**Proposed date of publication (dd/mm/yy):** April 2015

11.	<b>Proposed date of entry into force: <input checked="" type="checkbox"/> Six months from date of publication, and/or (dd/mm/yy):</b> <b><input type="checkbox"/> Trade facilitating measure</b>
12.	<b>Final date for comments: <input checked="" type="checkbox"/> Sixty days from the date of circulation of the notification and/or (dd/mm/yy):</b> 13 January 2015. Comments are welcomed only to MRLs that were changed (in bold in a document linked under point 5 of this notification). <b>Agency or authority designated to handle comments: <input checked="" type="checkbox"/> National Notification Authority, <input checked="" type="checkbox"/> National Enquiry Point. Address, fax number and e-mail address (if available) of other body:</b>  European Commission DG Health and Consumers, Unit G6-Multilateral International Relations Rue Froissart 101, B-1049 Brussels Tel: (+32 2) 295 42 63 Fax: (+32 2) 299 80 90 E-mail: sps@ec.europa.eu
13.	<b>Text(s) available from: <input checked="" type="checkbox"/> National Notification Authority, <input checked="" type="checkbox"/> National Enquiry Point. Address, fax number and e-mail address (if available) of other body:</b>  European Commission DG Health and Consumers, Unit G6-Multilateral International Relations Rue Froissart 101, B-1049 Brussels Tel: (+32 2) 295 42 63 Fax: (+32 2) 299 80 90 E-mail: sps@ec.europa.eu