NOTIFICATION

Addendum

The following communication, received on 12 April 2023, is being circulated at the request of the Delegation of the European Union.

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| Maximum residue levels for bromopropylate, chloridazon, fenpropimorph, imazaquin and tralkoxydim in or on certain products |
| The proposal notified in G/SPS/N/EU/527 (15 December 2021) is now adopted Commission Regulation (EU) 2023/710 of 30 March 2023 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 bromopropylate, chloridazon, fenpropimorph, imazaquin and tralkoxydim in or on certain products (Text with EEA relevance) [OJ L 93, 31 March 2023, p. 57–83].The MRLs for the active substance bifenthrin, which was initially included in the draft Regulation notified in G/SPS/N/EU/527, will be reviewed in a separated Regulation taking into account a recent assessment from the European Food Safety Authority (EFSA Journal 2023;21(3):7864). The proposal will be notified to the Committee on Sanitary and Phytosanitary Measures.The MRLs for the active substance fenpropimorph for barley, oat, rye, wheat, sugar beet roots, all tissues from mammals and milk, which were initially lowered to the limit of quantification in the draft Regulation notified in G/SPS/N/EU/527, are maintained at the level proposed by Codex Alimentarius in the adopted Commission Regulation (EU) 2023/710.The adopted Commission Regulation (EU) 2023/710 includes a provision to allow the marketing of products which have been produced in the Union or imported into the Union before the modified MRLs start applying.The Regulation shall apply from 21 October 2023.[https://members.wto.org/crnattachments/2023/SPS/EEC/23\_8914\_00\_e.pdf](https://members.wto.org/crnattachments/2023/SPS/EEC/23_8914_00_e.pdf%22%20%5Ct%20%22_blank)<https://members.wto.org/crnattachments/2023/SPS/EEC/23_8914_00_f.pdf><https://members.wto.org/crnattachments/2023/SPS/EEC/23_8914_00_s.pdf> |
| **This addendum concerns a:** |
| [ ] Modification of final date for comments |
| [**X**] Notification of adoption, publication or entry into force of regulation |
| [**X**] Modification of content and/or scope of previously notified draft regulation |
| [ ] Withdrawal of proposed regulation |
| [ ] Change in proposed date of adoption, publication or date of entry into force |
| [ ] Other:  |
| **Comment period: *(If the addendum extends the scope of the previously notified measure in terms of products and/or potentially affected Members, a new deadline for receipt of comments should be provided, normally of at least 60 calendar days. Under other circumstances, such as extension of originally announced final date for comments, the comment period provided in the addendum may vary.)*** |
| [ ] Sixty days from the date of circulation of the addendum to 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 A4-Multilateral International RelationsRue Froissart 101, B-1049 BrusselsTel: +(32 2) 295 4263Fax: +(32 2) 299 8090E-mail: sps@ec.europa.eu |
| **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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