NOTIFICATION

Addendum

The following communication, received on 6 May 2020, is being circulated at the request of the Delegation of the European Union.

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| Draft Commission Implementing Regulation concerning the non-renewal of the approval of the active substance mancozeb, in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending the Annex to Commission Implementing Regulation (EU) No 540/2011 |
| In the proposal notified in G/SPS/N/EU/384 (24 April 2020) the description of content should read:  This draft Commission Implementing Regulation provides that the approval of the active substance mancozeb is not renewed in accordance with Regulation (EC) No 1107/2009. Existing authorisations of plant protection products containing mancozeb will be withdrawn and such products can then no longer be placed on the market. The non-renewal of the approval is based on a scientific assessment conducted under Regulation (EC) No 1107/2009 by experts from the Member States of the European Union and the European Food Safety Authority (EFSA).  This decision only concerns the placing on the market of this substance and plant protection products containing it. Following non-approval and the expiry of all grace periods for stocks of products containing this substance, separate action will likely be taken on MRLs and a separate notification will be made in accordance with SPS procedures. |
| **This addendum concerns a:** |
| [ ] Modification of final date for comments |
| [ ] 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)*: |
| **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 Consumers, Unit G6-Multilateral International Relations  Rue Froissart 101, B-1049 Brussels  Tel: +(322) 295 4263  Fax: +(322) 299 8090  E-mail: [sps@ec.europa.eu](mailto: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:** |
| European Commission  DG Health and Consumers, Unit G6-Multilateral International Relations  Rue Froissart 101, B-1049 Brussels  Tel: +(322) 295 4263  Fax: +(322) 299 8090  E-mail: [sps@ec.europa.eu](mailto:sps@ec.europa.eu) |

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