

5 October 2022

(22-7504)

Committee on Sanitary and Phytosanitary Measures

## NOTIFICATION

Addendum

The following communication, received on 4 October 2022, is being circulated at the request of the Delegation of the <u>European Union</u>.

<u>Specific requirements for the performance of official controls on the use of pharmacologically active</u> <u>substances authorised as veterinary medicinal products or as feed additives and of prohibited or</u> <u>unauthorised pharmacologically active substances and residues thereof</u>

The proposal notified in G/SPS/N/EU/571 (20 June 2022) is now adopted by Commission Delegated Regulation (EU) 2022/1644 of 7 July 2022 supplementing Regulation (EU) 2017/625 of the European Parliament and of the Council with specific requirements for the performance of official controls on the use of pharmacologically active substances authorised as veterinary medicinal products or as feed additives and of prohibited or unauthorised pharmacologically active substances and residues thereof (Text with EEA relevance) [OJ L 248, 26 September 2022, p. 3].

This Regulation shall apply from 15 December 2022.

https://members.wto.org/crnattachments/2022/SPS/EEC/22 6763 00 e.pdf https://members.wto.org/crnattachments/2022/SPS/EEC/22 6763 00 f.pdf https://members.wto.org/crnattachments/2022/SPS/EEC/22 6763 00 s.pdf

## This addendum concerns a:

| []           | Modification of final date for comments                                      |
|--------------|--|
| [ <b>X</b> ] | Notification of adoption, publication or entry into force of regulation      |
| []           | 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 Commission DG Health and Food Safety, Unit D2-Multilateral International Relations Rue Froissart 101, B-1049 Brussels Original: English

Page: 1/2

- 2 -

Tel: +(32 2) 295 4263 Fax: +(32 2) 299 8090 E-mail: <u>sps@ec.europa.eu</u>

## 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 Food Safety, Unit D2-Multilateral International Relations Rue Froissart 101, B-1049 Brussels Tel: +(32 2) 295 4263 Fax: +(32 2) 299 8090 E-mail: <u>sps@ec.europa.eu</u>

\_\_\_\_\_