EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on *Veterinary Medicinal Products*
26 June 2023

CIRCABC Link: *https://circabc.europa.eu/ui/group/a2ba6dd7-f812-406a-8b1c-2aaf25ec4277/library/d00b8734-0f9a-4f13-90b3-95266d7f0ba1?p=1&n=10&sort=modified_DESC*

AGENDA

Section A Information and/or discussion

A.01 State of play of implementation of Regulation (EU) 2019/6 on Veterinary Medicinal Products.

A.02 Implementation of Regulation of the European Parliament and of the Council laying down transitional rules for the packaging and labelling of veterinary medicinal products authorised or registered in accordance with Directive 2001/82/EC and Regulation (EC) No 726/2004.

A.03 EMA Scientific advice under Article 107(6) of Regulation (EU) 2019/6 for the establishment of a list of antimicrobials which shall not be used in accordance with Articles 112, 113 and 114 of the same Regulation or which shall only be used in accordance with these articles subject to certain conditions - Presentation by EMA.

A.04 Information on next steps concerning the implementation of Commission Delegated Regulation (EU) 2023/905 supplementing Regulation (EU) 2019/6 of the European Parliament and of the Council as regards the application of the prohibition of use of certain antimicrobial medicinal products in animals or products of animal origin exported from third countries into the Union.

A.05 Exchange of views on the list of the abbreviations and pictograms common throughout the Union, in accordance with Article 17(2).

A.06 Exchange of views on on uniform rules on the size of small immediate packaging units, in accordance with Article 17(3).

Section B Draft(s) presented for an opinion

B.01 Exchange of views and technical agreement of the Committee on a draft Commission Implementing Regulation (EU) amending Regulation (EU) No 37/2010 to classify the substance **rafoxanide** as regards its maximum residue limit.

(PLAN/2023/1132)

Legal Basis: Regulation (EU) No 37/2010

Procedure: Examination procedure

B.02 Exchange of views and technical agreement of the Committee on a draft Commission Implementing Regulation (EU) amending Regulation (EU) No 37/2010 to classify the substance k**etoprofen** as regards its maximum residue limit.

(PLAN/2020/6400)

Legal Basis: Regulation (EU) No 37/2010

Procedure: Examination procedure