EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on *Veterinary Medicinal Products*
27 November 2024

CIRCABC Link: *https://circabc.europa.eu/ui/group/a2ba6dd7-f812-406a-8b1c-2aaf25ec4277/library/3326c8c3-132b-4532-ab2a-ee313aac54c6?p=1&n=10&sort=modified_DESC*

AGENDA

Section A Information and/or discussion

A.01 Discussion on a draft Implementing act on the establishment under Article 115(5) of Regulation (EU) 2019/6 of a list of substances which are essential for the treatment of equine species, or which bring added clinical benefit compared to other treatment options available for equine species and for which the withdrawal period for equine species shall be six months.

A.02 Development of implementing acts on GMP for veterinary medicinal products under Article 93(2) of Regulation (EU) 2019/6: status quo and next steps.

A.03 Discussion on the state of play on the implementation of Regulation (EU) 2022/839 of the European Parliament and of the Council (QRDv9 updates).

Section B Draft(s) presented for an opinion

B.01 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) amending Regulation (EU) 2021/17 establishing a list of variations not requiring assessment in accordance with Regulation (EU) 2019/6

(PLAN/2023/2278)

Legal Basis: Regulation (EU) 2019/6

Procedure: Examination procedure

Section C Draft(s) presented for discussion

C.01 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) amending Commission Regulation (EU) 2018/782 of 29 May 2018 establishing the methodological principles for the risk assessment and risk management recommendations referred to in Regulation (EC) No 470/2009

(PLAN/2024/1244)

Legal Basis: Regulation (EC) No 470/2009

Procedure: Examination procedure

C.02 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) amending Regulation (EU) No 37/2010 as regards chemical-unlike biological substances

(PLAN/2024/1245)

Legal Basis: Regulation (EC) No 470/2009

Procedure: Examination procedure

C.03 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) amending Implementing Regulation (EU) 2017/12 regarding the form and content of the applications and requests for the establishment of maximum residue limits in accordance with Regulation (EC) No 470/2009 of the European Parliament and of the Council

(PLAN/2024/2136)

Legal Basis: Regulation (EC) No 470/2009

Procedure: Examination procedure