EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on *Veterinary Medicinal Products*
31 January 2023

CIRCABC Link: *https://circabc.europa.eu/ui/group/a2ba6dd7-f812-406a-8b1c-2aaf25ec4277/library/15afb1f9-712a-4d42-ae16-b75ffc56c256*

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 Draft mandate to the European Medicines Agency for a scientific advice on the implementing measure under Article 115(5) of Regulation (EU) 2019/6 as regards the 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.03 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.

A.04 Information on the Commission Notice - Application of the Union's veterinary medicines acquis in markets historically dependent on medicines supply from or through parts of the United Kingdom other than Northern Ireland**.**

A.05 Update on the discussion as regards **Diclofenac** - Member State's update including any new information on reported vulture deaths in the Member States that have diclofenac authorised and potential new authorisations given at national level.

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) No 37/2010 to classify the substance **ketoprofen** as regards its maximum residue limit.

(PLAN/2020/6400)

Legal Basis: Regulation (EU) No 37/2010

Procedure: Examination procedure