



EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Veterinary Medicinal Products

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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** Collection of data on antimicrobial medicinal products used in animals (Article 57 of Regulation (EU) 2019/6) – Update from Member States.
- A.03** Recording of annual volume of sales in the Union product database (Article 58(12) Regulation (EU) 2019/6) – Update from Member States.
- A.04** Information on the legal framework under Regulation (EC) No 470/2009 and chemical-unlike biological substances.
- A.05** Discussion on a working document on the amendment of Regulation (EU) 2021/17 establishing a list of variations not requiring assessment in accordance with Regulation (EU) 2019/6.
- A.06** Discussion on a working document 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.07** Information on the discussions held at the first meeting of Expert Group on GMP for veterinary medicinal products under Article 93(2) of Regulation (EU) 2019/6.