EUROPEAN COMMISSION

Health and Food Safety Directorate General

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**Standing Committee on *Veterinary Medicinal Products***
**15 November 2023**

***CIRCABC Link:*** *https://circabc.europa.eu/ui/group/a2ba6dd7-f812-406a-8b1c-2aaf25ec4277/library/8f708794-8389-4c43-9f23-f650744e7b01?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** Discussion on the establishment under Article 107(6) of Regulation (EU) 2019/6 of a list of antimicrobials which shall not be used in accordance with Articles 112 and 113 of the same Regulation or which shall only be used in accordance with these articles subject to certain conditions

**A.03** Information on the state of play on 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.04** Collection of data on antimicrobial medicinal products used in animals – State of play on the implementation of Article 57 of Regulation (EU) 2019/6 at Member State level

**A.05** Recording of annual volume of sales in the Union product database – State of play on the implementation of Article 58(12)

**A.06** Information on ongoing files linked to MRLs:

- Information on a proposal for an amendment of Commission Regulation (EU) No 37/2010 as regards the application of MRL to dried, diluted, processed or compound foods
- Information on a proposal for an amendment of Commission Delegated Regulation (EU) 2019/2090 as regards a change in the definition of “unauthorised substance”

## Section B Draft(s) presented for an opinion

**B.01** Exchange of views and a technical agreement of the Committee on a draft Commission Implementing Regulation (EU) amending Regulation (EU) No 37/2010 as regards the classification of the substance **sodium salicylate** with respect to its maximum residue limit in foodstuffs of animal origin

(PLAN/2023/2348)

**Legal Basis:** Regulation (EU) No 37/2010

**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) adopting a list of the abbreviations and pictograms common throughout the Union to be used for the purposes of Articles 10(2) and 11(3) of Regulation (EU) 2019/6 of the European Parliament and of the Council of 11 December 2018 on veterinary medicinal products and repealing Directive 2001/82/EC

(PLAN/2023/921)

**Legal Basis:** Regulation (EU) 2019/6

**Procedure:** Examination procedure

**C.02** Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) adopting uniform rules on the size of small immediate packaging units referred to in Article 12 of Regulation (EU) 2019/6 of the European Parliament and of the Council of 11 December 2018 on veterinary medicinal products and repealing Directive 2001/82/EC

(PLAN/2023/922)

**Legal Basis:** Regulation (EU) 2019/6

**Procedure:** Examination procedure