SCHEDULE 5

Medicated feedingstuffs and specified feed additives

Sampling and analysis E+W+S

22.—(1) If any enforcement action is taken under this Schedule based on a sample, that sample must have been taken and analysed in accordance with **[**[F1]Regulation (EC) No 152/2009 laying down methods of sampling and analysis for the official control of feedingstuffs.**]**

(2) Unless otherwise specified in the marketing authorisation, it is a defence if the active ingredient in the medicated feedingstuff sample is within the following tolerances—

[[F2]Tolerance table for medicated feedingstuff**]**

Level of active ingredient specified on the label	*Tolerance*
≤500mg/kg	±30%
>500mg/kg ≤5g/kg	±20%
>5g/kg	±10%**]**

(3) Unless otherwise specified in the Commission Regulation authorising the specified feed additive in question, it is a defence if the active ingredient of a specified feed additive in a feedingstuff sample is within the tolerances set out in Articles 11(5) and paragraph 2(e) of Annex IV to Regulation (EC) No 767/2009 of the European Parliament and of the Council.

Extent Information

E1 This version of this provision extends to England and Wales and Scotland only; a separate version has been created for Northern Ireland only

Textual Amendments

F1 Words in Sch. 5 para. 22(1) substituted (26.3.2019) by The Veterinary Medicines and Animals and Animal Products (Examination of Residues and Maximum Residue Limits) (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/676), regs. 1(2)(a), **2(4)(e)**

F2 Sch. 5 para. 22(2) Table substituted (E.W.S.) (17.5.2024) by The Veterinary Medicines (Amendment etc.) Regulations 2024 (S.I. 2024/567), regs. 1(1), **143** (with reg. 207)

Sampling and analysis N.I.

22.—(1) If any enforcement action is taken under this Schedule based on a sample, that sample must have been taken and analysed in accordance with **[**[F3]Regulation (EC) No 152/2009 laying down methods of sampling and analysis for the official control of feedingstuffs.**]**

(2) Unless otherwise specified in the marketing authorisation, it is a defence if the active ingredient in the medicated feedingstuff sample is within the following tolerances—

Tolerance table for medicated feedingstuff

Level of active ingredient specified on the label	*Tolerance*
≤50 mg/kg	50%

Level of active ingredient specified on the label	*Tolerance*
>50 mg/kg ≤ 500 mg/kg	40%
>500 mg/kg ≤ 5g/kg	30%
>5g/kg ≤50g/kg	20%
>50g/kg	10%

(3) Unless otherwise specified in the Commission Regulation authorising the specified feed additive in question, it is a defence if the active ingredient of a specified feed additive in a feedingstuff sample is within the tolerances set out in Articles 11(5) and paragraph 2(e) of Annex IV to Regulation (EC) No 767/2009 of the European Parliament and of the Council.

Extent Information

E2 This version of this provision extends to Northern Ireland only; a separate version has been created for England and Wales and Scotland only

Textual Amendments

F3 Words in Sch. 5 para. 22(1) substituted (26.3.2019) by The Veterinary Medicines and Animals and Animal Products (Examination of Residues and Maximum Residue Limits) (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/676), regs. 1(2)(a), **2(4)(e)**

Status:
There are multiple versions of this provision on screen. These apply to different geographical extents.
Skip to:

– E+W+S - England, Wales and Scotland extent
– N.I. - Northern Ireland extent

Changes to legislation:
There are currently no known outstanding effects for the The Veterinary Medicines Regulations 2013, Paragraph 22.