Veterinary Medicines Directorate

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Post Authorisation Assessments

Tribex 50 mg/ml Oral Suspension for Sheep

Vm 08749/3146

22 July 2026	One-off alignment of the product information with version 9.1 of the QRD template.
06 January 2026	Submission of a Ph. Eur. CEP for an active substance.
19 September 2025	Change in legal entity of the Marketing Authorisation Holder from Chanelle Animal Health Ltd, 7 Rodney Street, Liverpool, L1 9EF, United Kingdom to Chanelle Pharmaceuticals Manufacturing Ltd, Loughrea, Co. Galway, H62 FH90, Ireland.
04 May 2024	Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile. (NI)
04 May 2024	Changes to the quality part of the dossier, deletion of a manufacturing site for an active substance. (NI)
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04 May 2024	Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile. (GB)
04 May 2024	Changes to the quality part of the dossier, deletion of a manufacturing site for an active substance. (GB)
26 October 2021	To change the mock ups design for the product.
26 July 2018	Updates to the SPC and product literature following the outcome of a periodic safety update report.
27 April 2017	Addition of a manufacturer of the active substance
20 July 2016	Change in the specification parameters of the active substance.
19 June 2015	Addition of a new manufacturing site for the active ingredient.
29 July 2013	Variation to update a test procedure of a related substance of the active substance.
13 May 2013	Grouped variation concerning: the deletion of a non-significant in-process test used during manufacture of the finished product, the extension of the finished product shelf life, and the widening of shelf life specifications.
07 May 2013	Variation to update the SPC and Labelling following the outcome of a European Commission Decision Article 35 Referral for all Veterinary Medicinal Products containing active substances belonging to the class of Flukicides.
20 September 2012	Grouped variation concerning: the deletion of an active substance manufacturer, a change in the name of the active substance manufacturer, the addition of an active substance manufacturer, and a change in the melting point specification of an active substance.

08 June 2011	Implementation of an additional Warning statement on the SPC/Labelling.
23 November 2010	Addition of an active substance manufacturer.
08 August 2008	EU Renewal.
02 July 2004	Addition of a user warning.