Veterinary Medicines Directorate

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Post Authorisation Assessments

Flukiver 50 mg/ml Oral Suspension

Vm 52127/3001

28 January 2026	One-off alignment of the product information with version 3 of the QRD template. Change to the meat and offal withdrawal period for sheep.
16 September 2025	Addition of test procedure for an active substance.
29 April 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
12 March 2025	Change of Marketing Authorisation Holder from: Elanco Europe Ltd., Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, United Kingdom to: Elanco GmbH, Heinz-Lohmann Strasse 4, Groden, D-27472 Cuxhaven, Germany.
05 September 2023	Minor change to a footnote on the shelf-life specification.
17 August 2023	Minor editorial changes to part 2 of the dossier. Minor editorial changes to part 2 of the dossier.
21 July 2023	Submission of an updated certificate of suitability.
02 December 2021	Minor changes to an approved test procedure of the finished product. Minor changes to the approved finished product shelf-life specifications.
15 July 2021	Change in the address of a manufacturer of the finished product, also responsible for batch release.
18 March 2021	Update of a test procedure to comply with the updated Ph. Eur. monograph.
01 October 2020	Change of Marketing Authorisation Holder from Eli Lilly and Company Limited, Lilly House, Priestley Road, Basingstoke, Hampshire, RG24 9NL to Elanco Europe Ltd, Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, United Kingdom.
26 June 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer
01 November 2019	Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Deletion of Ph. Eur. certificates of suitability for an active substance.
05 June 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
18 March 2014	Submission of a new Ph. Eur. Certificate of Suitability for a new manufacturer of the active and submission of an updated Ph. Eur. Certificate of Suitability for an already approved manufacturer.

28 February 2013	Updates to the SPC and Product Literature to bring in line with flukicide legislation
30 January 2013	Submission of an updated Ph. Eur. Certificate of Suitability for an active substance from an already approved manufacturer
24 October 2012	Change of MAH
07 March 2012	Change of distributor
28 July 2010	Change of specification of excipients to comply with Ph. Eur.
07 July 2010	Change of wording of storage condition of the active substance
02 June 2010	Updates to the SPC and Product Literature regarding mixing with other products
11 November 2009	Introduction of a retest period (36 months) and storage conditions for the active substance
27 October 2009	Submission of a new Ph. Eur. Certificate of Suitability for an active substance
17 December 2008	Change of legal category from PML to POM-VPS Changes to the SPC and Product Literature to bring in line with new legislation
05 June 2008	Change of shape of closure nozzle
07 March 2008	Change of address of the MAH
30 August 2006	Changes in test procedure for a starting material used in the manufacturing process of the active substance Change to specification of a reagent used in the manufacture of the active substance Change to the specification of the active substance Change to the specification of an intermediate produced during the manufacturing process of the active substance
15 August 2006	Change in test procedures for a starting material used in the manufacturing process of the active substance
12 July 2006	Renewal
24 January 2006	Change of manufacturing site of the finished product, quality control testing and batch release Change to specification of the finished product Change of composition of immediate packaging
21 November 2003	Changes to the manufacturing process of the active substance
12 September 2003	Renewal
24 August 2001	Addition of a manufacturing site of secondary assembly of the dosage form
16 December 1997	Renewal
09 December 1996	Change of specification of the active substance
10 January 1996	Change of manufacturing site of the dosage form
07 September 1995	Change of importer
16 May 1995	Change of withdrawal period for Meat from Sheep from 28 days to 42 days