Veterinary Medicines Directorate

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Post Authorisation Assessments

Switch 40 mg/ml Pour-on Solution

Vm 18182/5001

02 September 2026	One-off alignment of the product information with version 3 of the GB QRD templates.
17 March 2022	Update to user safety warnings.
22 March 2019	Change in the colour specification of the finished product.
04 July 2018	Change in storage conditions of the finished product.
14 February 2018	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
21 December 2016	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
27 January 2015	Deletion of a manufacturing site. Updates to the product packaging. Change in distributor details.
14 February 2012	Variation to change the name of the manufacturer and assembler of dosage form.
30 March 2011	Variation to change the name of a distributor.
30 March 2011	Variation to change the name of the Marketing Authorisation Holder.
15 June 2010	Variation to change the name of the active substance manufacturer.
15 June 2010	Variation to make minor changes to the manufacturing process for the finished product.
14 April 2010	Addition of a manufacturer.
26 March 2010	Addition of a manufacturer responsible for batch release.
13 March 2008	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from GSL to AVM-GSL.
19 December 2007	Variation to change the assembler of dosage form.
29 November 2007	Variation to change the address of the Marketing Authorisation Holder and distributor.
08 November 2005	Renewal.
19 October 2000	Renewal.
09 September 1999	Change of assembler of dosage form.
14 January 1999	Change of the name and address of the Marketing Authorisation Holder.