Veterinary Medicines Directorate

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Post Authorisation Assessments

Switch 4% w/v Pour-on Solution

Vm 18182/4000

•	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.