Veterinary Medicines Directorate

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Post Authorisation Assessments

Killitch 25% w/v Cutaneous Emulsion

Vm 01974/4003

13 May 2025	Change in the pharmacovigilance system master file location.
02 June 2023	Change in the name of the Marketing Authorisation Holder from Carr & Day & Martin Limited to Carr & Day & Martin.
30 September 2022	Replacement of a batch control testing site for a finished product.
04 May 2022	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
15 March 2022	Replacement of a manufacturing site of the finished product. Replacement of a primary packaging site of the finished product. Replacement of secondary packaging site of the finished product. Replacement of a manufacturing site of the finished product.
09 February 2022	Change in the address of the MAH from Carr & Day & Martin Limited, Animal House, Boundary Road, Lytham, Lancashire, FY8 5LT to Carr & Day & Martin Limited, Woodland Granaries, Narrow Lane, Wymeswold, Loughborough, LE12 6SD.
08 February 2022	Replacement of a manufacturer responsible for batch release of the finished product.
23 October 2020	Addition of a new container for the finished product.
16 October 2020	Deletion of a pack size of the finished product.
06 March 2020	Change in the name and address of the marketing authorisation holder from Carr Day and Martin Limited, Docklands, Dock Road, Lytham, Lancashire, FY8 5AQ to Carr & Day & Martin Limited, Animal House, Boundary Road, Lytham, Lancashire, FY8 5LT.
06 June 2019	Addition of a new specification parameter to the specification with its corresponding test method of the finished product Change in the specification limits of the finished product Changes in the composition (excipients) of the finished product Change to the batch formulation
27 March 2018	Change in the specification limits 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.
14 February 2012	Change in the name of the Manufacturer/Assembler of Dosage Form.
17 June 2008	Variation to change product distributor.
26 February 2008	Variation to bring the SPC/Labelling in line with Veterinary Regulations, 2005. Transfer of legal category from GSL to ASM-GSL.
20 December 2007	Variation to change the Manufacturer/Assembler of Dosage Form.
07 December 2007	Change in the Marketing Authorisation Holder.
16 November 2006	Change in Batch Size.
16 November 2006	Variation to change the Manufacturer/Assembler of Dosage Form.
02 November 2006	Addition of a QC Testing and Batch Release Site.
21 December 2005	Renewal.
28 November 2002	Change in pack details.
28 Novemeber 2002	Change in pack details.
29 May 2002	Change in composition of finished product. Change in shelf-life of finished product.
03 May 2002	Change of the Manufacturer of the finished product.
03 May 2002	Change of the Assembler of the finished product.
11 December 2001	Change in the name and address of the Marketing Authorisation Holder.
08 June 2001	Renewal.
24 June 1995	Renewal.