Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Killitch 25% w/v Cutaneous Emulsion

Vm 01974/4003

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| • | 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.<br>Replacement of a primary packaging site of the finished product.<br>Replacement of secondary packaging site of the finished product.<br>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<br>Change in the specification limits of the finished product<br>Changes in the composition (excipients) of the finished product<br>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. |

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| • | 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. |