

Post Authorisation Assessments

Vectin Horse Oral Paste 18.7 mg/g Vm 50146/3006

08 April 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
19 January 2025	Minor change in the manufacturing process of the finished product.
23 April 2024	Deletion of a CEP for the manufacture of an active substance. (NI) Deletion of a CEP for the manufacture of an active substance. (NI)
23 April 2024	Submission of a new Ph. Eur. CEP from a new manufacturer for a non-sterile active substance. (NI)
23 April 2024	Deletion of a CEP for the manufacture of an active substance. (GB) Deletion of a CEP for the manufacture of an active substance. (GB)
13 April 2024	Submission of a new Ph. Eur. CEP from a new manufacturer (replacement or addition) for a non-sterile active substance. (NI)
28 September 2021	Change in the fill volume of the finished product.
30 March 2021	Minor changes in the SPC, whereby amendments are made to typographical errors in Section 4.5 and 4.7 of the SPC.
18 March 2021	Replacement of a secondary packaging site of the finished product.
26 January 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
27 July 2020	Change in distributor details from Intervet UK Ltd, Walton Manor, Walton, Milton Keynes, MK7 7AJ to MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ.
25 September 2019	Change in the invented name of the veterinary medicinal product from Diapec P Gel to Bimectin Paste in Germany only.
15 August 2019	Change in the name address of the manufacturer of the finished product. Change in the name and address of a manufacturer of the finished product, also responsible for batch release. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
18 October 2018	Change of MAH, from Cross Vetpharm Group Ltd., Broomhill Road, Tallaght, Dublin 24, Ireland to Bimeda Animal Health Limited, 2 / 3 / 4 Airton Close, Tallaght,

	Dublin 24, Ireland.
12 March 2018	Change in RMS from UK to IE.
12 January 2016	Submission of an updated certificate of suitability.
18 October 2012	Submission of a new European Pharmacopoeia Certificate of Suitability for a new active substance manufacturer.
29 March 2012	Grouped variation concerning the submission of updated European Pharmacopoeia Certificates of Suitability for already approved active substance manufacturers.
03 October 2008	Renewal (UK as RMS).
19 September 2008	Submission of an updated European Pharmacopoeia Certificate of Suitability for an already approved active substance manufacturer.
20 July 2006	Extension of the finished product shelf life.
16 December 2005	Addition of a site for assembly only.
22 September 2005	Change of distributor.
17 August 2005	Variation to harmonise the SPC between the UK and IE.
17 August 2005	Repeat use procedure.
23 July 2004	Variation to change the name of the veterinary medicinal product in PT only.
09 October 2003	New EUDE (UK as RMS)
19 June 2002	New Marketing Authorisation.