



Post Authorisation Assessments

Bimeprazol 370 mg/g Oral Paste for Horses Vm 50146/4043

26 March 2026	One-off alignment of the product information with version 3 of the QRD template.
08 September 2025	Submission of a Ph. Eur. CEP for an active substance.
05 September 2025	Change in distributor.
11 January 2025	Deletion of a non-significant in-process test during the manufacture of the finished product.
02 October 2024	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance.
02 October 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
28 April 2024	Minor changes:- to an approved test procedure for the finished product.
22 November 2022	Updated certificate of suitability from an already approved manufacturer.
09 August 2022	Transfer the method of the release test for the finished product.
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.
06 November 2020	Changes to the labelling and package leaflet.
08 October 2020	Change of MAH from Oy Medfiles Ltd, Volttikatu 5, Kuopio, FI-70700, Finland to Bimeda Animal Health Limited, 2, 3, 4 Airton Close, Tallaght, Dublin 24, IE-D24 E032, Ireland.
06 October 2020	Change in distributor details. From Oy Medfiles Ltd, Volttikatu 5, Kuopio, FI-70700, Finland to Cross Vetpharm Group UK Ltd, Unit 2, Bryn Cefni, Llangefni, Anglesey, LL77 7XA.