

## **Post Authorisation Assessments**

### **Vetroxy LA 200 mg/ml Solution for Injection for Cattle, Sheep and Pigs** Vm 50146/4021

18 February 2025	Alignment of the product information with version 9.0* of the QRD templates.
29 February 2024	Change in site of manufacture, primary packaging and analytical testing. Change in site of batch release. Change in site of secondary packaging batch release
12 July 2022	Submission of a new or updated Ph. Eur. certificate of suitability.
22 December 2021	Renewal - UK as CMS.
15 April 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
18 March 2021	Replacement of a secondary packaging site of the finished product.
23 September 2019	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.
19 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.
15 March 2018	Change in RMS from UK to NL.
12 December 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.