



## Post Authorisation Assessments

### Carprox vet 50 mg/ml Solution for Injection for Dogs and Cats

Vm 01656/4014

•	01 February 2023	Change to comply with an update of the relevant monograph of the Ph. Eur. Reflecting compliance with the Ph. Eur. by removing reference to the internal test method and test method number for the active substance.
•	07 November 2019	Change in the manufacturer of a starting material used in the manufacturing process of the active substance.
•	11 June 2019	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	10 May 2017	Minor change to the restricted part of an Active Substance Master File.
•	23 December 2015	Minor changes to an approved test procedure Minor changes to an approved test procedure Deletion of a manufacturing site. Deletion of a manufacturing site.
•	23 December 2015	Renewal - UK as CMS
•	16 May 2014	Changes to the manufacturing process for the active substance.
•	28 February 2014	Change in the manufacturing process of the finished product.
•	02 October 2013	Change in test procedure for an excipient.  Two changes to comply with Ph. Eur and in-house specifications.  Change the specification parameters of an excipient.
•	26 September 2013	Change of distributor
•	19 September 2013	Addition of a manufacturing site and site of primary packaging
•	16 September 2013	Change of supplier of packaging components
•	20 June 2013	Change in name of product from 'Rycarfa 50mg Tablets for Dogs' to 'Carprox vet 50mg Tablets for Dogs'
•	11 February 2013	To update the ASMF for an already approved ASM.
•	23 <sup>rd</sup> November 2012	Extension of the finished product shelf life from 2 years to 3 years.
•	21 April 2011	Repeat Use Comm – UK as CMS

