



## Post Authorisation Assessments

### Buprevet Multidose 0.3 mg/ml Solution for Injection for Dogs and Cats Vm 57446/4003

•	07 July 2023	Change in name of manufacturer of the finished product.
•	09 June 2023	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
•	23 January 2023	Change in MAH from Richter Pharma AG, Feldgasse 19, 4600 Wels, Austria to VetViva Richter GmbH, Durisolstrasse 14, 4600 Wels, Austria.
•	17 January 2023	Updated certificate of suitability from an already approved manufacturer.
•	04 November 2022	Updated certificate of suitability from an already approved manufacturer.
•	26 April 2021	Increase in batch size (400 L) of the finished product.
•	18 June 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	28 November 2018	Renewal – UK as CMS.
•	15 May 2018	Minor changes to an approved test procedure of the finished product.
•	23 February 2017	Change in section 4.6 of the SPC.
•	23 February 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	26 June 2015	Change in the batch size of the finished product.
•	01 April 2015	Deletion of a Ph. Eur. Certificate of Suitability.
•	19 December 2014	Submission of an updated Ph. Eur. Certificate of Suitability for the active substance.