



Post Authorisation Assessments

Ketavet 100 mg/ml Solution for Injection for Dogs, Cats and Horses Vm 42058/3003

•	08 October 2024	Change in any part of the primary packaging material not in contact with the finished product formulation.
•	08 October 2024	Minor change to an approved test procedure for the finished product.
•	01 July 2024	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
•	28 April 2024	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
•	03 August 2023	One-off alignment of the product information with version 9.0 of the QRD template.
•	05 May 2020	Increase in the shelf-life of the finished product as packaged for sale, from 2 years to 3 years.
•	03 February 2020	Renewal - UK as CMS
•	12 November 2019	Change in the address of the marketing authorisation holder from Zoetis UK Limited 5th Floor, 6 St. Andrew Street London EC4A 3AE to Zoetis UK Limited 1st Floor, Birchwood Building Springfield Drive Leatherhead Surrey KT22 7LP.
•	25 September 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	17 April 2018	Change in RMS from UK to ES.
•	29 December 2016	Change in the address of the marketing authorisation holder in France, Czech Republic & Slovakia.
•	05 April 2016	Changes to the manufacturing process.
•	05 May 2015	Change to the QPPV contact details.