



Post Authorisation Assessments

Euthasol Vet. 400 mg/ml, Solution for Injection Vm 19994/4016

22 February 2026	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible. (NI)
30 September 2025	Alignment of the product information with version 9.0* of the QRD templates.
07 May 2025	Change in the immediate packaging of the finished product.
03 July 2024	Increase in the batch size of a finished product.
28 April 2024	Minor change to the approved test procedure for finished product. (NI)
13 April 2024	Minor change to the approved test procedure for finished product. (GB)
16 November 2023	Minor changes to an approved test procedure for the finished product. (GB)
07 April 2022	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
05 November 2019	Introduction of a new pharmacovigilance system.
25 March 2019	Change in RMS from UK to IE.
22 March 2019	Change of distributor to Dechra Veterinary Products Limited, Sansaw Business Park, Hadnall, Shrewsbury, Shropshire SY4 4AS, United Kingdom
17 July 2018	Changes to the labelling and/or package leaflet.
22 December 2016	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
20 October 2016	Renewal – UK as RMS
26 June 2013	Approval of Unforeseen Mock Ups
14 November 2012	Repeat Use Procedure.