



Post Authorisation Assessments

Rimadyl Cattle 50 mg/ml Solution for Injection Vm 42058/5152

12 February 2026	Minor change to the restricted part of an active substance master file.
16 October 2025	Change in the specification parameters or limits of the finished product to describe more accurately the appearance of the product.
20 May 2025	Alignment of the product information with version 3.0* of the QRD templates.
26 January 2025	Minor changes: – to an approved test procedure for active substance.
26 January 2025	Changes in the manufacturing process of the active substance: - Minor change to the restricted part of an Active Substance Master File.
06 June 2024	Substantial changes in the updated version of the ASMF of Carprofen.
14 July 2023	Deletion of a non-significant specification parameter in the specification parameters or limits of an excipient. Deletion of a non-significant specification parameter in the specification parameters or limits of an excipient. Deletion of a non-significant specification parameter in the specification parameters or limits of an excipient. Deletion of a non-significant specification parameter in the specification parameters or limits of an excipient. Deletion of a non-significant specification parameter in the specification parameters or limits of an excipient.
06 July 2023	Deletion of a non-significant specification parameter of an active substance.
06 July 2023	Change to comply with Ph. Eur. or with a national pharmacopoeia of a Member State.
08 March 2023	Addition of a secondary packaging site. Addition of a secondary packaging site.
03 November 2022	Change in the name of a quality control testing site. Deletion of a manufacturing site for the active substance.
12 October 2022	Addition of a secondary packaging site of a finished product. Addition of a secondary packaging site of a finished product.
03 December 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.

27 September 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
06 April 2016	A change to the restricted part of the ASMF.
01 May 2015	Changes to the QPPV contact details.
29 April 2015	Addition of an active substance manufacturer.
17 January 2014	Changes to composition of finished product. Changes to the composition of the stopper. Change to pack size of finished product – addition of 100 ml and 250 ml sizes.
09 October 2013	Change in the name and address of the Marketing Authorisation Holder in AT, BE, FR, LU only from Pfizer to Zoetis.
03 October 2013	Changes to an existing pharmacovigilance system as described in the DDPS.
16 August 2013	Change in distributor and MAH from Pfizer Limited to Zoetis UK Limited.
22 February 2013	Renewal – Ireland as RMS.
9 February 2012	Change in the name of the address of the MAH in Spain only.
15 December 2009	Replacement of the manufacturing site of the finished product.
13 November 2009	Alignment of specifications of active substance with new Ph. Eur monograph for carprofen for veterinary use.