



Post Authorisation Assessments

Rimadyl Palatable Tablets 50 mg for Dogs

Vm 60021/3015

18 February 2026	Addition of a quality testing site for the finished product.
09 January 2026	Submission of a new Ph. Eur. CEP for a new manufacturer of the active substance.
19 December 2025	Adverse events section: Footnote 4 has been removed in order to maintain alignment on packaging with IE.
22 October 2025	Minor changes to an approved test procedure for an excipient in the finished product.
25 September 2025	Alignment of the product information with version 9.0* of the QRD templates.
09 September 2025	Minor changes: — to an approved test procedure for a finished product.
11 March 2025	Change in the qualitative composition of the immediate packaging.
14 November 2024	Change in legal entity of MA holder for UK(NI) from Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP to Zoetis Belgium S.A., 2nd Floor, Building 10, Cherrywood Business Park, Loughlinstown, Co. Dublin, Ireland.
August 2023	Deletion of a manufacturing site for an active substance.
29 March 2022	Change in shape or dimensions of the container or closure (immediate packaging).
24 March 2022	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product for solid pharmaceutical forms.
24 March 2022	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product for solid pharmaceutical forms.
09 April 2020	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, First Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP. Change(s) in the SPC, Labelling or Package Leaflet of veterinary medicinal product intended to implement the outcome of a procedure concerning PSUR.
June 2018	Change in the specification limits of the finished product.
06 April 2016	A change to the restricted part of the ASMF.
30 March 2016	Deletion of a manufacturing site (for an active substance, intermediate or finished product, packaging site, manufacturer responsible for batch release, site where batch control takes place, or supplier of a starting material, reagent or excipient (when mentioned in the dossier)).

20 October 2015	Change in name of manufacturer.
09 June 15	Change in dimensions of the immediate packaging container.
06 December 2013	Addition of an alternative API manufacturer.
14 February 2013	Variation to delete an active substance manufacturer.
15 April 2009	Variation to align the product specifications with the new European Pharmacopoeia Monograph.
14 February 2008	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of legal category from POM to POM-V.
09 January 2008	Renewal.
27 March 2006	Change in the specification of an excipient.
23 February 2006	Batch Control.
07 February 2006	Change in the storage conditions of the finished product.
02 February 2006	Addition of a site for secondary assembly.
11 January 2006	Variation concerning the 'Indications'.
04 November 2005	Batch Control.
27 October 2005	Batch Control.
27 October 2005	Batch Control.
27 September 2005	Change to the packaging shape and dimensions.
23 June 2005	Addition of a distributor.
17 March 2005	Changes to the finished product.
23 February 2004	Change in the specifications of an excipient to comply with the European Pharmacopoeia.
30 September 2003	Variation to change the name of the active substance manufacturer.