Veterinary Medicines Directorate

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Post Authorisation Assessments

Rimadyl Palatable Tablets 100 mg for Dogs

Vm 60021/3013

31 July 2026	Change to importer, batch control arrangements and quality testing (replacement or addition of a site) for a finished product.
16 June 2026	Change in source of an excipient or reagent with TSE risk.
13 March 2026	Other changes to the active substance: - Substantial changes in the updated version of the ASMF.
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.
01 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.
19 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).
26 February 2016	Change in dimensions of the immediate packaging container.
18 November 2015	Change in the name of manufacturer for the finished product.
20 October 2015	Change in name of manufacturer.
09 June 2015	Change in dimensions of the immediate packaging container.
06 December 2013	Addition of an alternative API manufacturer.
14 February 2013	Deletion of an active substance manufacturer.
15 April 2009	Variation to align the product specifications with the new European Pharmacopoeial Monograph.
14 February 2008	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
09 January 2008	Renewal.
21 July 2006	Batch Control.
27 March 2006	Change in the specification of an excipient.
07 February 2006	Change in the storage conditions of the finished product.
02 February 2006	Addition of a site of secondary assembly.
11 January 2006	Variation concerning a change to the ‘Indications’ of the product.
27 September 2005	Variation to increase the container shape and dimensions.
23 June 2005	Addition of a distributor.
17 March 2005	Changes to a test procedure for the finished product.
23 February 2004	Change in an excipients specifications to comply with European Pharmacopoeia.
30 September 2003	Change in the name of the active substance manufacturer.