

Post Authorisation Assessments

Rimadyl Palatable Tablets 20 mg for Dogs

Vm 60021/3014

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| 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. |
| 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 |

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| | place, or supplier of a starting material, reagent or excipient (when mentioned in the dossier). |
| 18 November 2015 | Change in the name of manufacturer for the finished product. |
| 20 October 2015 | Change in name of manufacturer. |
| 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 | Renewal. |
| 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 September 2005 | Change to the packaging shape and dimensions. |
| 24 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. |