Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Rimadyl Palatable Tablets for Dogs 100 mg

Vm 42058/4120

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| • | 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.<br>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. |

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| • | 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. |