



## Post Authorisation Assessments

### Rycarfa 50 mg/ml Solution for Injection for Dogs and Cats Vm 01656/4174

|                  |   |
|------------------|---|
| 22 January 2025  | One-off alignment of the product information with version 9.0* of the QRD templates.  |
| 11 June 2024     | Change in the manufacturing process of the finished product.  |
| 18 May 2024      | Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.  |
| 01 February 2023 | Change to comply with an update of the relevant monograph of the Ph. Eur.<br>Reflecting compliance with the Ph. Eur. by removing reference to the internal test method and test method number for the active substance. |
| 06 November 2019 | Change in the manufacturer of a starting material used in the manufacturing process of the active substance.  |
| 11 June 2019     | Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.   |
| 13 February 2019 | Renewal - UK as CMS.  |
| 07 June 2018     | Deletion of manufacturing site for an active substance.<br>Deletion of manufacturing site for an active substance.<br>Deletion of manufacturing site for an active substance.   |
| 09 May 2017      | Minor change to the restricted part of an Active Substance Master File.   |
| 16 March 2016    | Minor changes to an approved test procedure<br>Minor changes to an approved test procedure  |
| 19 May 2015      | Addition of UK local representative information to the package leaflet.   |