Veterinary Medicines Directorate

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Post Authorisation Assessments

Rycarfa 50 mg/ml Solution for Injection for Dogs and Cats

Vm 01656/4174

•	01 February 2023	Change to comply with an update of the relevant monograph of the Ph. Eur. 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. Deletion of manufacturing site for an active substance. 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 Minor changes to an approved test procedure
•	19 May 2015	Addition of UK local representative information to the package leaflet.