



Post Authorisation Assessments

Rilexine 75 mg Tablets for Dogs and Cats

Vm 05653/4131

02 July 2025	Change in the specification parameters and/or limits of the finished product Change in the specification parameters and/or limits of the finished product Change to in-process tests or limits applied during the manufacture of the finished product. Change to in-process tests or limits applied during the manufacture of the finished product.
24 June 2025	One-off alignment of the product information with version 3 of the QRD template.
09 February 2023	Change in test procedure for an excipient.
21 December 2022	Change in the shelf-life or storage conditions of the finished product.
07 October 2022	Change in the specification parameters and/or limits of the finished product.
17 May 2022	Deletion of a non-significant in-process test applied during the manufacture of the finished product. Change in the specification parameters and/or limits of the immediate packaging of the finished product. Change in the specification parameters and/or limits of the immediate packaging of the finished product.
31 July 2020	Changes to the labelling and/or package leaflet.
10 January 2020	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
23 January 2018	Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State. Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State
08 September 2017	Update of the test procedure to comply with the updated general Ph. Eur monograph. Increase in the shelf-life of the finished product as packaged for sale, from 24 to 36 months.
06 October 2016	Harmonisation of SPC and mock-ups with IE.
26 November 2015	Change in the specification parameters of the finished product. Deletion of a non-significant specification parameter.
19 March 2013	Addition of a new active substance manufacturer

	with a European Pharmacopoeial Certificate of Suitability.
26 October 2012	Grouped variation to change a test procedure, reduce the impurity limits of a test procedure, and to change the sample number of a test procedure.
07 April 2011	Renewal.
09 January 2009	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
21 August 2008	Addition of a new pack size.
27 February 2008	Extension of indications.
04 October 2006	Minor alteration to the SPC following advice from the VMD.