



## Post Authorisation Assessments

For details of post authorisation assessments prior to 1<sup>st</sup> January 2021,  
please refer to the [EMA](#) website.

### **Felisecto Plus 30 mg/5 mg spot-on solution for cats >2.5–5 kg** Vm 42058/5025

• May 2024	Alternate test method for a starting material added.
• May 2024	One-off alignment of the product information with version 9.0 of the QRD templates.
• 11 May 2024	Change in name of manufacturer of starter material. Change in addresses for manufacturers of a starter material. Manufacturing site of a starting material deleted. Manufacturing site for a starting material deleted.
• 04 May 2024	Addition of a new specification parameter for a starting material.
• 18 April 2024	Unlimited renewal.
• 20 October 2023	Extension of the re-test period of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier.
• 04 August 2023	Change in batch size of active substance. Change in batch size of active substance. Change in batch size of active substance.
• 18 April 2023	Addition of an alternative supplier of a starting material.
• April 2023	Change in batch size of an intermediate used in the manufacturing process of the active substance.
• April 2023	Change in batch size of an intermediate used in the manufacturing process of the active substance.
• April 2023	Change in address of supplier of the active substance.
• April 2023	Change in name and address of a manufacturer of the active substance.
• April 2023	Change in the name of a supplier of starting material.
• April 2023	Changes to a test procedure for the immediate packaging of the active substance. Change in manufacturer of the active substance.
• April 2023	Update the Summary of Product Characteristics and product literature with regard to adverse reactions, and to update the product literature text in line with the current template.