



Post Authorisation Assessments

Tramazole 100 mg/ml Oral Suspension for Cattle and Sheep Vm 11810/4011

04 December 2025	One-off alignment of the product information with version 3 of the QRD template
10 October 2025	Submission of a Ph. Eur. CEP for an active substance.
30 April 2025	Changes in the composition (excipients) of the finished product.
15 May 2024	Change of name of the veterinary medicinal product from Tramazole 100 mg/ml SC Oral Suspension for Cattle and Sheep to Tramazole 100 mg/ml Oral Suspension for Cattle and Sheep.
07 June 2023	Change in the shelf-life of the finished product. Change in the shelf-life or storage conditions of the finished product.
27 October 2021	National renewal
05 May 2021	Removal of manufacture sites. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
17 June 2020	Change in the specification limits of the finished product. Change in the specification limits of the finished product. Widening of the in-process limits applied during the manufacture of the finished product.
05 April 2018	Increase in the shelf-life of the finished product from 2 to 3 years.
12 April 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance manufacturer.