



Post Authorisation Assessments

Vetoryl 20 mg Chewable Tablet for Dogs Vm 50406/5005

05 January 2026	One-off alignment of the product information with version 3.0* of the QRD templates.
23 September 2025	Addition of a new specification parameter to the specification with its corresponding test method. Addition of a new specification parameter to the specification with its corresponding test method. Addition of a new specification parameter to the specification with its corresponding test method. Deletion of a test procedure for the active substance. Deletion of a non-significant specification parameter of an active substance. Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
23 September 2025	Submission of a new Ph. Eur. certificate of suitability for a starting material used in the manufacturing of the active substance.
10 September 2025	To delete a non-significant specification parameter from the release and shelf-life specification of the finished product.
10 September 2025	To widen the disintegration time in-process limits, applied during the manufacture of the finished product.
08 May 2025	Change in the name of the veterinary medicinal product from Vetoryl Flavoured 20 mg Tablets for Dogs, to Vetoryl 20 mg Chewable Tablet for Dogs.
07 January 2025	Addition of a Local Representative for reporting suspected adverse events- Dechra Veterinary Products Limited, Sansaw Business Park, Hadnall, Shrewsbury, Shropshire, SY4 4AS, United Kingdom, Tel: +44 (0) 1939 211200.