



Post Authorisation Assessments

Tialin 125 mg/ml Solution for Use in Drinking Water for Pigs, Chickens and Turkeys Vm 50406/3036

05 August 2025	Change in the manufacturing process of the finished product.
19 May 2025	Change of Marketing Authorisation Holder from Dechra Limited, Snaygill Industrial Estate, Keighley Road, Skipton, North Yorkshire, BD23 2RW, United Kingdom to Dechra Regulatory B.V., Handelsweg 25, 5531 AE Bladel, The Netherlands.
19 December 2023	Alignment with version 9 of the QRD templates.
26 October 2022	Unlimited renewal.
02 March 2022	Change in shape or dimensions of the container or closure (immediate packaging). Changes in the qualitative and quantitative composition of the immediate packaging of the finished product. Changes in the qualitative and quantitative composition of the immediate packaging of the finished product.
11 February 2022	Deletion of manufacturer responsible for batch release and finished product.
30 March 2021	Deletion of manufacturing site for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
21 May 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
12 February 2019	Changes to an existing pharmacovigilance system as described in the DDPS.
25 September 2018	Addition of a manufacturer responsible for batch release including batch control/testing. Addition of a secondary packaging site of the finished product. Addition of a primary packaging site of the finished product. Addition of a manufacturing site of the finished product.
30 August 2018	Change in RMS from UK to NL.