



## Post Authorisation Assessments

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### Tulaxa 100 mg/ml Solution for Injection for Cattle, Pigs and Sheep Vm 01656/5073

14 April 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
18 May 2024	One-off alignment of the product information with version the latest version of the QRD templates.
04 July 2023	Change in test procedure for the finished product to comply with Ph. Eur. (NI)
16 June 2023	Unlimited renewal
14 April 2023	Change in test procedure for the finished product to comply with Ph. Eur.
20 October 2021	Change in the SPC, labelling or package leaflet following assessment of the same change for the reference product.
17 February 2021	Tightening of specification limits of an active substance used in the manufacturing process of the active substance.
19 August 2020	Minor changes to an approved test procedure of the finished product
24 September 2019	Increase in the shelf-life of the finished product as packaged for sale, from 24 months to 36 months.