Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

Trymox LA 150 mg/ml Suspension for Injection for Cattle, Sheep, Pigs, Dogs, Cats

Vm 05150/5003

3 August 2026	Change in the finished product specification parameters. Change in the finished product specification parameters. Change in a finished product test procedure. Change in a finished product test procedure.
13 July 2026	Submission of a new CEP for the manufacture of an active substance. Introduction of a re-test period for an active substance.
31 March 2026	Submission of a new Ph. Eur. certificate of suitability for a manufacturer of the active substance. Introduction of a re-test period/storage period supported by real time data.
22 February 2026	Submission of a Ph. Eur. CEP for an active substance.
16 December 2025	Submission of mock ups for joint labelling with IE.
25 November 2024	One-off alignment of the product information with version 9.0 of the QRD templates.
17 April 2024	Submission of an updated CEP for amoxicillin trihydrate.
30 December 2022	Updated certificate of suitability from an already approved manufacturer.
10 August 2022	Change in Distributor, from: Tullyvin Laboratories Ltd, 35 Abbeydale Park, Newtownards, Co. Down, Northern Ireland, BT23 8RE to: FORTE Healthcare Limited, Block 3, Unit 9, CityNorth Business Campus, Stamullen, Co. Meath, K32 D990, Republic of Ireland.
15 February 2021	Changes to the SPC/product labelling/package leaflet following an Article 35 referral.
22 May 2020	Deletion of manufacturing site for an active substance. Qualitative and/or quantitative changes to the excipients.
22 May 2020	Increase in the shelf-life of the finished product as packaged for sale, from 18 months to 24 months (50 & 100 ml) and 12 months to 24 months (250ml).
30 August 2019	Change in the shelf-life of the finished product as packaged for sale.