Veterinary Medicines Directorate

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Post Authorisation Assessments

Tribovax T

•	06 December 2018	Replacement of a manufacturer responsible for batch release of the finished product. Addition of a manufacturer responsible for batch release of the finished product.
•	02 June 2010	Submission of a new European Pharmacopoeia Certificate of Suitability for an active substance manufacturer.
•	07 May 2010	Renewal.
•	08 April 2010	Variation to change the Marketing Authorisation Holder.
•	12 December 2007	Variation to change a finished product test procedure.
•	16 August 2007	Variation to replace an in-process test procedure.
•	04 April 2007	Variation to update the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from PML to POM-VPS.
•	19 October 2005	Reviewed MA.