Veterinary Medicines Directorate

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Post Authorisation Assessments

Equip T

Vm 42058/4064

•	30 September 2022	Change in name of the manufacturer of the active substance.
•	22 July 2022	Removal of the non-toxicity and in vivo residual toxicity test from in-process testing. Replacement of the absence of toxin and irreversibility of toxoid test with the absence of tetanus toxoid test performed on the purified toxoid bulk.
•	19 August 2020	Change in the address of the marketing authorisation holder from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
•	09 April 2020	Changes to a test procedure for the finished product.
•	19 November 2018	Deletion of manufacturing site for an active substance.
•	18 November 2015	Addition of a new presentation form
•	13 December 2011	Addition of a dosing device
•	14 November 2011	Renewal
•	14 September 2011	Change to manufacturing process of the finished product Change to test procedure performed on the finished product
•	30 December 2010	Change of name of manufacturing site for blending, filling and assembly, QC testing labelling and batch release
•	01 July 2010	Addition of a manufacturing site for secondary packaging
•	29 June 2010	Addition of a manufacturing site of batch release
•	21 April 2009	Change of MAH
•	10 September 2008	Replacement of a test performed on the finished product
•	21 August 2008	Harmonisation of the SPC
•	16 August 2007	Replacement of an in process test performed on the finished product
•	04 April 2007	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in line with new legislation