



Post Authorisation Assessments

Equip T Suspension for Injection

Vm 42058/5232

04 August 2025	One-off alignment of the product information with version 3.0 of the GB National SPC/QRD templates.
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