



Post Authorisation Assessments

Equip F Vm 42058/5231

14 October 2025	Submission of mock ups.
04 March 2025	G.I.18 update of the product information to version 3 of the National template.
14 June 2024	a. The addition of acceptable countries of origin as source for the porcine pancreas used to make the porcine trypsin powder. b. The removal of the countries of origin for the lactose used in the production process of porcine trypsin. c. To make some editorial changes in the TSE risk assessment.
25 March 2024	Change in the source of a starting material used in the manufacturing process of the active substance.
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.
03 December 2019	Changes to a test procedure for the starting material.
26 October 2018	Changes to a test procedure for the active substance
18 November 2015	Addition of a new presentation form
23 April 2013	Change of manufacturer of a starting material used in the manufacture of the active substance
13 December 2011	Addition of administration device
14 November 2011	Renewal
14 September 2011	Addition of manufacturing site responsible for control testing and full batch release of finished product
30 December 2010	Change of name of manufacturer site for the active substance, 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 for batch release
21 April 2009	Change of MAH
21 August 2008	Harmonisation of SPC
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