



Post Authorisation Assessments

Stabox 5% w/w Premix for Medicated Feeding Stuff for Piglets Vm 05653/4044

26 January 2026	Submission of mock ups.
26 November 2025	One-off alignment of the product information with version 3 of the GB QRD templates.
07 March 2025	Deletion of a manufacturing site for an active substance. Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance.
21 October 2024	Change in the address of a manufacturer of the finished product (including batch release or quality control testing sites).
21 December 2023	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance.
12 January 2022	Addition of a new specification parameter to the specification with its corresponding test method of the finished product. Tightening of specification limits of the finished product. Replacement to a test procedure for the finished product. Replacement to a test procedure for the finished product. Change in storage conditions of the finished product. Change in the specification parameters and/or limits of the finished product. Change in the specification parameters and/or limits of the finished product.
24 November 2021	Change in the name of a manufacturer of the finished product, also responsible for batch release.
11 December 2020	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
27 June 2019	Submission of a new Ph. Eur. certificate of suitability for an active substance. Introduction of a re-test period of the active substance.
06 June 2018	Change to part of the (primary) packaging material not in contact with the finished product formulation.
02 September 2009	Variation to bring the SPC/ Labelling in line with the Veterinary Regulations, 2005. Transfer of legal category from POM to POM-V.
05 March 2008	Addition of an active substance manufacturer.
05 March 2008	Addition of an active substance manufacturer.
19 December 2007	Variation to change a site of manufacture.
29 November 2005	Renewal.
24 October 2002	Renewal.
16 December 1999	Variation to change the address of a distributor.

02 November 1995

New Marketing Authorisation.