Veterinary Medicines Directorate

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Post Authorisation Assessments

Vanguard 7
Vm 42058/4157

•	25 March 2024	Change in the source of a starting material used in the manufacturing process of the active substance.
•	18 November 2022	Introduction of in vitro antigen capture ELISA potency test. Use of antigen capture ELISA to quantify antigen.
•	06 March 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. Change in the SPC, Labelling or Package Leaflet of veterinary medicinal products intended to implement the outcome of a procedure concerning PSUR. Update SPC/QRD in line with the latest QRD template.
•	19 October 2018	Change in the specification parameters and/or limits of the finished product.
•	09 November 2016	Change in the composition (excipients) of the finished product.
•	19 February 2014	Transfer of MA from Pfizer Ltd to Zoetis UK Limited and change of distributor. Change of name of the active substance manufacturer and change of name of the finished product manufacturer responsible for batch release.
•	16 December 2010	Renewal.
•	18 November 2010	Removal of a test.
•	07 July 2010	Change in the onset of immunity period for one of the active substances.
•	17 February 2010	Addition of suppliers of starting materials.
•	15 July 2009	Change to the wording of Section 4.2 of the SPC.
•	13 February 2009	Change to an active substance strain designation.
•	24 May 2006	Changes to the SPC and product literature to bring them into line with new legislation.
•	24 May 2006	Change to legal category from POM to POM-V.