Veterinary Medicines Directorate

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Post Authorisation Assessments

Covexin 8

Vm 42058/4023

•	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.
•	30 April 2019	Changes in the manufacturing process of the active substance.
•	01 March 2018	Submission of a new Ph. Eur. TSE certificate of suitability for a starting material from a new or already approved manufacturer.
•	04 July 2017	Change in test procedure for active substance or starting material/reagent/intermediate used in the manufacturing process of the active substance
•	03 August 2016	Deletion of non-significant in-process tests.
•	14 April 2014	Transfer of MA from Pfizer Ltd to Zoetis UK Limited, change of distributor and change of name of the finished product manufacturer responsible for batch release.
•	21 November 2012	Replacement of manufacturing site for blending, filling and assembly
•	24 July 2012	Replacement of manufacturing site for secondary packaging Replacement of manufacturing site for batch release
•	18 January 2012	Changes to an existing pharmacovigilance system as described in the DDPS
•	02 September 2010	Renewal
•	02 April 2009	Change of MAH Change of Distributor
•	12 December 2007	Change in test procedure performed on the finished product
•	16 August 2007	Change of in-process test performed on the finished product
•	04 April 2007	Change of legal category from PML to POM-VPS Changes to the SPC and Product Literature to bring in line with new legislation