Veterinary Medicines Directorate

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Post Authorisation Assessments

Blackleg Vaccine
Vm 60021/3004

•	25 November 2024	Change in legal entity of MA holder for NI from Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP to Zoetis Belgium S.A., 2nd Floor, Building 10, Cherrywood Business Park, Loughlinstown, Co. Dublin, Ireland.
•	28 September 2022	Change in name and address of the manufacturer 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.
•	14 February 2017	Change in test procedure for the finished product.
•	13 June 2016	Deletion of a non-significant in-process test applied during the manufacture of the active substance Deletion of a non-significant in-process test applied during the manufacture of the active substance Deletion of a non-significant in-process test applied during the manufacture of the active substance Deletion of a non-significant in-process test applied during the manufacture of the active substance Deletion of an in-process test applied during the manufacture of the active substance.
•	21 March 2014	Transfer of MA from Pfizer Ltd to Zoetis UK Limited as well as change of distributor and change to the name of the finished product manufacturer.
•	25 September 2013	Change of manufacturing site for the active substance and in process control testing. Change in manufacturing process of the active substance and change in test procedure performed on the active substance. Change in immediate packaging of the active substance.
•	06 February 2013	Change in test procedure performed on the active substance
•	24 July 2012	Change of manufacturer of the finished product, secondary packaging and manufacturer responsible for quality control and batch release.
•	19 December 2011	Changes to an existing pharmacovigilance system as described in the DDPS.
•	02 September 2010	Renewal.
•	02 April 2009	Change of MA holder from Schering-Plough Ltd to Pfizer Ltd.

•	12 December 2007	Change of test procedure performed on the finished product.
•	04 April 2007	Changes to bring the SPC and Product Literature in line with new legislation and to change of legal category from PML to POM-V.
•	19 October 2005	Review.