Veterinary Medicines Directorate

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Post Authorisation Assessments

Deosect 5% w/v Concentrate for Cutaneous Spray Solution

Vm 42058/4033

•	26 August 2020	Change in the address of the MAH 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.
•	24 October 2019	Change in the specification limits of the finished product.
•	03 September 2018	Change in the manufacturer of a starting material of the active.
•	13 December 2017	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product. Replacement of a specification parameter of the finished product.
•	12 February 2016	Change in immediate packaging of the finished product
•	26 March 2014	Transfer of MA from Pfizer Ltd to Zoetis UK Limited, change of distributor and change to the name of the finished product manufacturer responsible for batch release.
•	18 January 2012	Changes to an existing pharmacovigilance system as described in the DDPS
•	23 March 2011	Change of manufacturer of the finished product
•	03 March 2011	Change of manufacturer for assembly of the finished product Change of manufacturing site for batch release and quality control testing of the finished product
•	09 February 2011	Change of manufacturer of the active substance
•	23 February 2010	Change of MAH
•	31 July 2008	Change of legal category from PML to POM-VPS Changes to the SPC and Product Literature to bring in line with new legislation
•	30 October 2007	Renewal
•	24 January 2007	Corrections to the SPC and Product Literature
•	29 November 2004	Renewal
•	05 November 2004	Changes to the SPC to bring in line with new legislation
•	25 June 2004	Deletion of indication for poultry
•	06 April 2004	Change of MAH
•	16 October 2001	Change of address of manufacturer of the active substance
•	20 July 2001	Change of product name from ‘Deosan Deosect’ to ‘Deosect Spray’
•	25 May 2001	Change of manufacturing site of assembly of the dosage form

•	16 January 2000	Change of distributor
•	16 June 1998	Change of manufacturing site for assembly of the dosage form
•	28 May 1998	Renewal
•	13 March 1997	Change of MAH