Veterinary Medicines Directorate

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Post Authorisation Assessments

Canac Flea and Tick Spot-on Solution 742 mg

Vm 16516/4083

•	04 May 2023	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
•	23 November 2022	Change in the specification parameters and/or limits of an excipient.
•	17 May 2019	Change in the invented name of the veterinary medicinal product from Beaphar Dog Flea and Tick Drops, 65% w/w, cutaneous solution' to 'Canac Flea and Tick Spot-on solution 742 mg'.
•	11 March 2019	Change of specification(s) of a former non Pharmacopoeial active substance to comply with the Ph. Eur. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	21 September 2017	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product.
•	09 August 2017	Change in the specification limits of the finished product
•	02 April 2013	Change in shape/dimensions of packaging Change of supplier of packaging components
•	22 February 2013	Change of supplier of an excipient
•	29 December 2011	Change in name of the manufacturer of the active substance
•	08 October 2008	Addition of supplier of packaging components
•	11 September 2008	Change in shape/dimensions of container/closure system
•	20 May 2008	Renewal
•	29 April 2008	Changes to the SPC and Product Literature to bring in line with new legislation
•	15 November 2007	Change of product name from ‘Canac Dog Flea and Tick Drops’ to ‘Beaphar Dog Flea and Tick Drops’
•	26 September 2007	Change of legal category from GSL to AVM-GSL
•	31 May 2007	Change of manufacturer of the active substance
•	06 July 2005	Renewal
•	18 December 2003	Addition of safety warnings regarding toxicity to cats
•	16 May 2003	Addition of packaging presentation Change of product name from ‘Friends Dog Flea Drops with Permethrin’ to ‘Canac Dog Flea and Tick Drops’
•	15 March 2002	Addition of a warning to the SPC and Product Literature
•	30 July 2001	Addition of manufacturer of the dosage form
•	11 July 2001	Change of MAH