Veterinary Medicines Directorate

United Kingdom
Veterinary Medicines Directorate
Woodham Lane
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NATIONAL PROCEDURE

PUBLICLY AVAILABLE ASSESSMENT REPORT FOR A VETERINARY MEDICINAL PRODUCT

Canac Dog Flea and Tick Collar, 15% w/w (Dimpylate)

Date Created: June 2018

MODULE 1

PRODUCT SUMMARY	01935/2017
Name, strength and pharmaceutical form	Canac Dog Flea and Tick Collar, 15% w/w (Dimpylate)
Applicant	Beaphar Ltd Rook Tree Farm Withersfield Road Great Wratting Suffolk CB9 7HD United Kingdom
Active substance	Dimpylate (as dimpylate stabilised)
ATC Vetcode	QP53AF03
Target species	Dogs
Indication for use	Treatment of dogs infested with fleas (*Ctenocephalides felis* and *C. canis*) and ticks (*Ixodes ricinus*).

MODULE 2

The Summary of Product Characteristics (SPC) for this product is available on the Product Information Database of the Veterinary Medicines Directorate.

(www.gov.uk/check-animal-medicine-licensed)

MODULE 3

PUBLIC ASSESSMENT REPORT

Legal basis of original application	Informed Consent application in accordance with Article 13c of Directive 2001/82/EC as amended.
Date of conclusion of the procedure	26th April 2018

I. SCIENTIFIC OVERVIEW

The quality, safety and efficacy aspects of this product are identical to Beaphar Flea and Tick Collar for Dogs, 15% w/w. The initial application for Beaphar Flea and Tick Collar for Dogs, 15% w/w was assessed before there was a requirement to have a public assessment report; therefore no details in this section are available.

V OVERALL CONCLUSION AND BENEFIT– RISK ASSESSMENT

The data submitted in the dossier demonstrate that when the product is used in accordance with the Summary of Product Characteristics the benefit/risk profile of the product is favourable.

MODULE 4

POST-AUTHORISATION ASSESSMENTS

The SPC and package leaflet may be updated to include new information on the quality, safety and efficacy of the veterinary medicinal product. The current SPC is available on the Product Information Database of the Veterinary Medicines Directorate website.

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The post-authorisation assessment (PAA) contains information on significant changes which have been made after the original procedure which are important for the quality, safety or efficacy of the product.

The PAA for this product is available on the Product Information Database of the Veterinary Medicines Directorate website.

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