Veterinary Medicines Directorate

United Kingdom
Veterinary Medicines Directorate
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NATIONAL PROCEDURE

PUBLICLY AVAILABLE ASSESSMENT REPORT FOR A VETERINARY MEDICINAL PRODUCT

Johnson's Flea Collar for Cats, 15% w/w.

Date Created: 01/12/2017

MODULE 1

PRODUCT SUMMARY

Name, strength and pharmaceutical form	Johnson's Flea Collar for Cats, 15% w/w
Applicant	Beaphar (UK) Ltd Rook Tree Farm Withersfield Road Great Wratting Suffolk CB9 7HD
Active substance	Dimpylate (as dimpylate stabilised)
ATC Vetcode	QP53AF03
Target species	Cats
Indication for use	Treatment of cats infested with fleas (*Ctenocephalides felis*)

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)

As the product was refused a marketing authorisation, a Summary of Product Characteristics was not approved for this product,

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	21/09/2017

I. SCIENTIFIC OVERVIEW

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

II. 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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