ASSURING THE SAFETY, QUALITY AND EFFICACY OF VETERINARY MEDICINES

United Kingdom
Veterinary Medicines Directorate
Woodham Lane
New Haw
Addlestone
Surrey KT15 3LS

NATIONAL PROCEDURE

PUBLICLY AVAILABLE ASSESSMENT REPORT FOR A VETERINARY MEDICINAL PRODUCT

MV Chlorhexidine (0.425% w/w) RTU Teat Dip and Teat Spray Solution

MODULE 1

PRODUCT SUMMARY

Name, strength and pharmaceutical form	MV Chlorhexidine (0.425% w/w) RTU Teat Dip and Teat Spray Solution
Applicant	Diversey Limited Weston Favell Centre Northampton Northamptonshire NN3 8PD
Active substance(s)	Chlorhexidine Gluconate solution
ATC Vetcode	QG52A
Target species	Bovine animals (cows)
Indication for use	A ready to use teat dip / spray for use as an aid in the control of bovine mastitis in cattle

MODULE 2

The Summary of Product Characteristics (SPC) for this product is available on the Veterinary Medicines Directorate website (www.vmd.defra.gov.uk)

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.

I. SCIENTIFIC OVERVIEW

The quality / safety / efficacy aspects of this product is/are identical to Deosan Teatcare Plus 0.425% w/w Teat Dip and Teat Spray Emulsion. The initial application for Deosan Teatcare Plus 0.425% w/w Teat Dip and Teat Spray Emulsion 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 for the target species is favourable and the quality and safety of the product for humans and the environment is acceptable.

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.

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

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.

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