



**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**

**Thixoshield 40 mg/g Teat Dip Solution**

**Date Created: December 2015**

## **MODULE 1**

### **PRODUCT SUMMARY**

Name, strength and pharmaceutical form	Thixoshield 40 mg/g Teat Dip Solution
Applicant	Diversey Limited Weston Favell Centre Northampton Northamptonshire NN3 8PD
Active substance	Benzyl alcohol
ATC Vetcode	QG52A
Target species	Cattle – milking cows
Indication for use	A ready to use teat dip as an aid in the control of bovine mastitis.

## **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](http://www.gov.uk/check-animal-medicine-licensed)

## **MODULE 3**

### **PUBLIC ASSESSMENT REPORT**

Legal basis of original application	Informed Consent application in accordance with Article 13 (c) of Directive 2001/82/EC as amended
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#### **I. SCIENTIFIC OVERVIEW**

The quality / safety / efficacy aspects of this product are identical to Deosan Thixodip BA, 4.0% w/w Teat Dip Solution. The initial application for Deosan Thixodip BA, 4.0% w/w Teat Dip Solution 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(s) 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.

([www.gov.uk/check-animal-medicine-licensed](http://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](http://www.gov.uk/check-animal-medicine-licensed))