

## **SUMMARY OF PRODUCTS CHARACTERISTICS**

### **1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Star Iodocare Concentrate Teat Dip and Teat Spray Solution 2.00%w/v

### **2. QUALITATIVE AND QUANTITATIVE COMPOSITION**

<b><u>Active substances</u></b>	<b><u>%w/v</u></b>
Available iodine	2.00

#### **Other relevant constituents**

Glycerol	5.45
Sorbitol Solution ( 70% w/w )	16.35

For a full list of excipients, see section 6.1

### **3. PHARMACEUTICAL FORM**

Concentrate for Teat Dip/Teat Spray Solution  
A brown liquid.

### **4. CLINICAL PARTICULARS**

#### **4.1 Target species**

Cattle – milking cows

#### **4.2 Indications for use, specifying the target species**

A dilutable teat dip/spray as an aid in the control of bovine mastitis.

#### **4.3 Contraindications**

Not applicable

#### **4.4 Special warnings for each target species**

None

#### **4.5 Special precautions for use**

i. Special precautions for use in animals

Wash and dry udders and teats before milking.

Teat dip cups should be emptied after milking and washed before re-use.

These are both important aspects of good hygiene and mastitis control.

For external use only.

- ii. Special precautions for the person administering the veterinary medicinal product to animals

**CONCENTRATE**

The following safety phrases relate to the concentrated product only, and do not apply once the product is diluted to the working solution

Risk of serious damage to eyes. Wear eye/face protection when preparing the dip or spray solution

**DILUTED WORKING SOLUTION**

When used as a spray, avoid working in spray mist.

Avoid contact with eyes. In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.

If swallowed, seek medical advice immediately and show this label.

Hands and exposed skin should be washed after using the product.

Do not eat drink or smoke while using the product.

Keep away from food , drink and animal feedstuffs

- iii. Other precautions

none

**4.6 Adverse reactions (frequency and seriousness)**

Very rare - change of active ingredient teat dip type can on very rare occasions cause skin irritation.

**4.7 Use during pregnancy, lactation or lay**

Can be used during pregnancy and lactation.

**4.8 Interaction with other medicinal products and other forms of interaction**

The use of the product has no known interactions with other products, including cow diet supplements.

Do not mix with other chemicals.

**4.9 Amount(s) to be administered and administration route**

This product is diluted before use. Prepare a fresh solution daily.

Teat dipping:

Add 1 part product to 3 parts of clean water and mix well

Directly after milking each cow, dip the full length of each teat in the product.

The teat cup should be kept topped up as necessary.

Teat spraying:

Add 1 part product to 4 parts of clean water and mix well

Directly after milking each cow, spray the entire surface of each teat with the product.

**4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary**

Not applicable for the intended mode of application.

**4.11 Withdrawal period(s)**

Withdrawal period for meat/milk - zero days/hours.

**5. PHARMACOLOGICAL PROPERTIES**

**Pharmacotherapeutic group:** Antiseptics and disinfectants, Iodine products

**ATC Vet Code:** QD08AG01

Teat dips

**5.1 Pharmacodynamic properties**

Iodine based teat dips have broad spectrum antibacterial action against mastitis causative organisms. The microbiological action of iodine appears to be due to an oxidative – reductive reaction involving various cell wall constituents which are irreversibly transformed. It appears sulphhydryl linkages, in bacterial cell wall components are specifically targeted by the iodine.

**5.2 Pharmacokinetic properties**

The absorption of iodine through the skin from teat dipping applications is well below levels which would indicate pharmacokinetic activity of the type described in the committee for veterinary medicinal products summary report on iodine.

**5.3 Environmental properties**

Iodine based teat dips are harmful to fish and aquatic life. Ponds, watercourses or ditches must not be contaminated with the product or used containers. The impact of the active ingredient (iodine) entering the environment via normal use of the product is low

**6. PHARMACEUTICAL PARTICULARS**

**6.1 List of excipients**

Glycerol

Sorbitol Solution ( 70% w/w )

Alcohol ( C13-C15) 12 mole ethoxylate  
citric acid monohydrate  
sodium hydroxide  
sulphuric acid ( for pH adjustment )  
water, deionised

## **6.2 Incompatibilities**

Not to mix the product with other medicinal product.

## **6.3 Shelf life**

Shelf life of the veterinary medicinal product as packaged for sale: 18 months.

## **6.4 Special precautions for storage**

Do not store above 25°C.  
Protect from direct sunlight.  
Store in tightly closed original container.  
Protect from frost  
Discard any remaining diluted solutions at the end of the day

## **6.5 Nature and composition of immediate packaging**

5,25, and 200 litres natural or opaque , white or blue high density polyethylene drum with grey or white high density polyethylene or polypropylene cap ( screw fit )

20 litre, blue , white or natural opaque, high density polyethylene drums, with black high density polyethylene cap ( screw fit ) ( tamper evident ) with expanded polyethylene gasket

Not all pack sizes may be marketed.

## **6.6 Special precautions for the disposal of unused veterinary medicinal product or waste materials derived from the use of such products, if appropriate**

Any unused veterinary medicinal product or waste material derived from such veterinary medicinal products should be disposed of in accordance with local requirements. To dispose of unused product to land you must have an authorisation under the Groundwater regulations 1998.  
Harmful to fish and aquatic life. Do not contaminate ponds, waterways or ditches with the product or used container'

**7. MARKETING AUTHORISATION HOLDER**

Diversey Limited  
Weston Favell Centre  
Northampton  
Northamptonshire  
NN3 8PD

**8. MARKETING AUTHORISATION NUMBER(S)**

**Vm** 15985/4025

**9. DATE OF FIRST AUTHORISATION**

**Date:** 24/08/1998

**10. DATE OF REVISION OF THE TEXT**

**Date:** October 2011