LABEL

<PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE>

{ High-density polyethylene 5, 10, 20, 60 or 200 litre cans }

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Blockade 0.25% w/w iodine teat dip solution.

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Active substance:

Available iodine	0.25 % w/w
Equivalent to	2.5 mg/g
or	12.8 mg per 5 ml dose

A viscous red-brown liquid

3. PHARMACEUTICAL FORM

Teat dip solution.

4. PACKAGE SIZE

5, 10, 20, 60 or 200 litre. Not all pack sizes may be marketed.

5. TARGET SPECIES

(Dairy) cows.

6. INDICATION(S)

Teat disinfection as an aid in the prevention of mastitis.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Dip each teat of the cow immediately after milking in a dip cup containing undiluted product. Ensure that the teat is covered to three quarters of length and replenish the dip cup as necessary. Always clean the dip cup after its use. Dosage: 5 ml per cow per treatment.

8. WITHDRAWAL PERIOD

Meat and offal: zero days / milk : zero hours.

9. SPECIAL WARNING(S), IF NECESSARY

Special precautions for use in animals:
For external use only.
The use in injured teats may delay wound healing process. It is recommended to discontinue the treatment until the teats are cured.
Allow the product to dry before the cows are exposed to wet (rainy), cold or windy weather conditions.
Contraindications:
Do not use in cases of hypersensitivity to the active substance or to any of the excipients.
Interaction with other medicinal products and other forms of interaction:
The use of this product in the specified manner (topical antiseptic) has no known interactions with other products, animal's diet and feed supplements.
User warnings:
Persons with known hypersensitivity to iodine or to any of the excipients should avoid using the product. If you develop symptoms following exposure, such as skin rash you should seek medical advice and show the label to the physician. Avoid ingesting the product. In case of accidental ingestion seek medical advice immediately and show the label to the physician. Do not eat, drink or smoke while using the product. This product might be mildly irritating to skin and eyes. Avoid contact with skin and eyes when administrating the product. If the product comes into contact with the eyes, rinse immediately with plenty of water. Wash hands after use.
Incompatibilities:
In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.
Use during pregnancy, lactation or lay:
Indicated for use in lactating and pregnant cattle.
Adverse reactions:
If you notice any serious effects or other effects not mentioned in this label, please inform your veterinary surgeon.

10. EXPIRY DATE

Batch number and expiry date: see label top of can. Do not use this veterinary medicinal product after the expiry date which is stated on the top of the can after EXP. The expiry date refers to the last day of that month.

11. SPECIAL STORAGE CONDITIONS

Store upright and tightly closed in the original container. Protect from frost. If product has frozen, thaw in a warm room and shake well before using. Protect from light. Do not store above 30°C.

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements. The product should not enter water courses as this may be dangerous for fish and other aquatic organisms. The 200 liter container should not be returned for refilling.

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, if applicable

For animal treatment only.

14. THE WORDS "KEEP OUT OF THE REACH AND SIGHT OF CHILDREN"

Keep out of sight and reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

DeLaval NV, Industriepark-Drongen 10, 9031 Gent, Belgium
Manufacturer responsible for batch release: DeLaval NV, Industriepark-Drongen 10, 9031 Gent, Belgium

16. MARKETING AUTHORISATION NUMBER

Vm 17140/4003

17. MANUFACTURER'S BATCH NUMBER

Batch number and expiry date: see label top of can.

D. Austin

Approved: 23 March 2017