

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGING WHERE THERE IS NO PACKAGE LEAFLET, i.e. Combined label and package leaflet

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

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Bovidip 20mg/ml Concentrate for Teat Dip or Spray Solution.

2. COMPOSITION

Active substance:

2 g per 100 ml as available iodine (concentrate).
25 mg per 5 ml dose as available iodine (ready-to-use solution).

Excipients:

Glycerol, 100 mg/ml

Clear Brown Liquid.

3. PACKAGE SIZE

5, 10, 20, 60 or 200 litre.

4. TARGET SPECIES

Cattle (dairy cattle)

5. INDICATIONS FOR USE

Indications for use

Teat disinfection as an aid in the prevention of mastitis.

6. CONTRAINDICATIONS

Contraindications

Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

7. SPECIAL WARNINGS

Special warnings

Do not mix with other chemicals. Prior to milking, wash teats with an udder wash solution and dry with a disposable paper towel. Discard any product that becomes contaminated. Always use a clean spray or dip cup and clean after use.

Special precautions for safe use in the target species:

For external use only. Allow product to dry before exposing the cows to wet (rainy), cold or windy conditions. Use in injured teats may delay the wound-healing process. It is recommended that treatment be discontinued until teat lesions have resolved. If signs of disease persist or appear, consult a veterinary surgeon.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Care should be taken to avoid eye contact. In case of eye contact, flush the eyes with copious amounts of water and seek medical advice. In case of accidental ingestion, seek medical advice immediately and show the combined package leaflet and label to the physician. When used as a spray, avoid working in a spray mist. Wash hands after use. Persons with iodine allergy should wear gloves and mask. The use of gloves during milking and dipping or spraying is recommended to protect the skin and for hygienic milk collection.

Pregnancy and lactation:

Can be used during lactation and pregnancy.

Interactions 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 medicaments or nutrition.

Major incompatibilities:

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

8. ADVERSE EVENTS

Adverse events

None known.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed on this label, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder using the contact details on this label, or via your national reporting system. (Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>, e-mail: adverse.events@vmd.gov.uk).

9. DOSAGE FOR EACH TARGET SPECIES, ROUTES AND METHOD OF ADMINISTRATION

Dosage for each species, routes and method of administration

About 5ml of the diluted product per cow per application.

10. ADVICE ON CORRECT ADMINISTRATION

Advice on correct administration

Dilute before use. Prepare a fresh solution daily. Dilute one part of the veterinary medicinal product with three parts of clean water and mix well. Always clean the dip cup or spray container after use.

Administration route:

- Dipping: Dip each teat immediately after milking in a teat dip cup containing diluted product. Dip the full length of the teats and replenish the dip cup as necessary.
- Spraying: Spray the entire surface of the teats after each milking.

11. WITHDRAWAL PERIODS

Withdrawal periods

Meat and offal: zero days.

Milk: zero hours.

12. SPECIAL STORAGE PRECAUTIONS

Special storage precautions

Keep out of the sight and reach of children.

Store upright in the tightly closed original container. Do not store above 25°C. Protect from light. Protect from frost. If the product has frozen, thaw in a warm place and shake well before use. For the larger pack sizes, the product should be rolled sufficiently to mix the solution. Under no circumstances should an attempt be made to shake the 60 or 200 litre packs.

Do not use this veterinary medicinal product after the expiry date which is stated on this label after Exp. The expiry date refers to the last day of that month.

13. SPECIAL PRECAUTIONS FOR DISPOSAL

Special precautions for disposal

Medicines should not be disposed of via wastewater.

The veterinary medicinal product should not enter water courses as iodine may be dangerous for fish and other aquatic organisms. Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any national collection systems. These measures should help to protect the environment.

The 200L container should not be returned for re-filling.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

14. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Classification of veterinary medicinal products

Veterinary medicinal product not subject to prescription.

15. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Marketing authorisation number:

Vm 17140/4008

Pack sizes

5, 10, 20, 60 or 200 litres high-density polyethylene drums with screw closures and seals. Not all pack sizes may be marketed.

16. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

17. CONTACT DETAILS

Contact details

Marketing authorisation holder and manufacturer responsible for batch release and contact details to report suspected adverse events:

DeLaval NV, Industriepark-Drongen 10, 9031 Gent, Belgium. PHV phone number: 0032 9 351 24 27

18. OTHER INFORMATION

UK only: AVM-GSL

Date on which the label was last revised: 12/2024

19. THE WORDS "FOR ANIMAL TREATMENT ONLY"

For animal treatment only.

20. EXPIRY DATE

Batch number and expiry date: see label top of can.
Exp {mm/yyyy}

Shelf life after first opening the immediate packaging: 6 months.
Shelf life after dilution according to directions: 1 day.

21. BATCH NUMBER

Lot {number}

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

Gavin Hall
Approved: 22 August 2025