

Blockade 2.56 mg/ml teat dip solution

Authorised

- Iodine (125I)

Product identification

Medicine name:

Blockade 2.56 mg/ml teat dip solution

Active substance:

Iodine (125I)

Target species:

Cattle

Route of administration:

Teat use

Product details

Active substance and strength:

Iodine (125I)

2.56 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Teat dip solution

Withdrawal period by route of administration:**Teat use:**

-

Cattle

- Meat and offal. 0 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD08AG03

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Package description:

A dark liquid contained in 5 liters, grey high-density polyethylene drums with screw closures and seals

A dark liquid contained in 10 liters, grey high-density polyethylene drums with screw closures and seals

A dark liquid contained in 20 liters, grey high-density polyethylene drums with screw closures and seals

A dark liquid contained in 60 liters, grey high-density polyethylene drums with screw closures and seals

A dark liquid contained in 200 liters, grey high-density polyethylene drums with screw closures and seals

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Delaval

Marketing authorisation date:

30/06/2025

Manufacturing sites for batch release:

Delaval

Responsible authority:

Danish Medicines Agency

Authorisation number:

72988

Date of authorisation status change:

30/06/2025

Reference member state:

Ireland

Procedure number:

IE/V/0485/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Iceland Italy Latvia Lithuania
Luxembourg Netherlands Norway Poland Portugal Romania Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 24/08/2025

Updated on: 13/03/2026

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Combined File of all Documents

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