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Dermastitis-Blocker 3 mg/ml, teat dip solution for cattle

Not authorised

- Iodine

Product identification

Medicine name:
Dermastitis-Blocker 3 mg/ml, teat dip solution for cattle

Active substance:
Iodine

Target species:
Cattle

Route of administration:
Teat use

Product details

Active substance and strength:
Iodine
3.08 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Teat dip solution

Withdrawal period by route of administration:

Teat use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD08AG03

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
Denmark

Package description:
(ID6): 1 unspecified outer container with 1 Barrel (High Density PolyEthylene) with 194.7 litre(s) (194.7 litre(s))
(ID3): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 19.5 litre(s) (19.5 litre(s))
(ID2): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 9.7 litre(s) (9.7 litre(s))
(ID5): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 58.4 litre(s) (58.4 litre(s))
(ID7): 1 unspecified outer container with 1 Barrel (High Density PolyEthylene) with 973.2 litre(s) (973.2 litre(s))
(ID4): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 24.3 litre(s) (24.3 litre(s))
(ID1): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 4.9 litre(s) (4.9 litre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ferdinand Eimermacher GmbH und Co. KG

Marketing authorisation date:
4/05/2018

Manufacturing sites for batch release:
Ferdinand Eimermacher GmbH und Co. KG

Responsible authority:
Danish Medicines Agency

Authorisation number:
60464

Date of authorisation status change:
16/02/2026

Reference member state:
Germany

Procedure number:
DE/V/0180/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 16/02/2022
Updated on: 14/03/2026
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