

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 subject to veterinary prescription except for some pack sizes

Authorisation status:

Surrendered

Authorised in:

France

Package description:

(ID1): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 4.9 litre(s) (4.9 litre(s))

(ID6): 1 unspecified outer container with 1 Barrel (High Density PolyEthylene) with 194.7 litre(s) (194.7 litre(s))

(ID4): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 24.3 litre(s) (24.3 litre(s))

(ID7): 1 unspecified outer container with 1 Barrel (High Density PolyEthylene) with 973.2 litre(s) (973.2 litre(s))

(ID3): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 19.5 litre(s) (19.5 litre(s))

(ID5): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 58.4 litre(s) (58.4 litre(s))

(ID2): 1 unspecified outer container with 1 Canister (High Density PolyEthylene) with 9.7 litre(s) (9.7 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:

23/04/2018

Manufacturing sites for batch release:

Ferdinand Eimermacher GmbH und Co. KG

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/9166557 7/2018

Date of authorisation status change:

7/01/2026

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)

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