

DECTOSPOT 10 MG/ML POUR-ON SOLUTION FOR CATTLE AND SHEEP

Not
authorised

- Deltamethrin

Product identification

Medicine name:

DECTOSPOT 10 MG/ML POUR-ON SOLUTION FOR CATTLE AND SHEEP

Active substance:

Deltamethrin

Target species:

Cattle
Sheep

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Deltamethrin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Pour-on solution

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

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Cattle

- Meat and offal. 18 day
- Milk. 0 day

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Sheep

- Meat and offal. 35 day
 - Milk. 24 hour
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC11

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Latvia

Package description:

500 ml high density polyethylene flexipacks with twin neck dispenser, internal graduated calibration chamber and polypropylene heat-sealed screw cap.

1 litre high-density polyethylene flat bottom containers with polypropylene closures and tamper evident with induction heat-sealed wadding. A spouted cap is provided with the 1 litre presentations.

2.5 litre high-density polyethylene flat bottom containers with polypropylene closures and tamper evident with induction heat-sealed wadding. A spouted cap is provided with the 2.5 litre presentations.

250 ml high density polyethylene flexipacks with twin neck dispenser, internal graduated calibration chamber and polypropylene heat-sealed screw cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bimeda Animal Health Limited

Marketing authorisation date:

26/01/2016

Manufacturing sites for batch release:

Bimeda Animal Health Limited

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/16/0001

Date of authorisation status change:

21/01/2025

Reference member state:

France

Procedure number:

FR/V/0293/001

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.