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:

- **Cattle**
  - Meat and offal. 18 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 24 hour

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

English (PDF)
Published on: 2/04/2026
Download

eu-puar-frv0293001-mr-rpe210-en.pdf