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

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:
Valid

Authorised in:
France

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

500 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:
4/02/2016

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6739546 0/2016

Date of authorisation status change:
6/07/2021

Reference member state:
France

Procedure number:
FR/V/0293/001

Concerned member states:
Austria Belgium Denmark Estonia Finland Ireland Italy Latvia Lithuania Poland Portugal Romania Spain Sweden

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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

Combined File of all Documents

English (PDF)
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