# 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 Dectospot 10 mg/ml solução para unção contínua para bovinos e ovinos

#### **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:**

# 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:

Portugal

#### 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:

9/05/2016

# Manufacturing sites for batch release:

Bimeda Animal Health Limited

## **Responsible authority:**

Directorate General For Food And Veterinary

#### **Authorisation number:**

1010/01/16DFVPT

# Date of authorisation status change:

23/02/2024

#### **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**

Combined File of all Documents

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