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Spotinor 10 mg/ml Spot-on Solution for cattle and sheep

Authorised

- Deltamethrin

Product identification

Medicine name:
Spotinor 10 mg/ml Spot-on Solution for cattle and sheep

Active substance:
Deltamethrin

Target species:
Cattle
Sheep

Route of administration:
Spot-on use

Product details

Active substance and strength:
Deltamethrin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- **Cattle**
  - Meat and offal. 17 day
  - Milk. 0 day
- **Sheep**
  - Milk. no withdrawal period

Not authorised for use in ewes producing milk for human consumption.

- Meat and offal. 35 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC11

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID4) 2.5 litre(s): Box (Cardboard) with 1 Container (High Density PolyEthylene) with 2.5 litre(s), closed with Screw cap (PolyPropylene)
(ID3) 1 litre(s): Box (Cardboard) with 1 Container (High Density PolyEthylene) with 1 litre(s), closed with Screw cap (PolyPropylene)
(ID2) 500 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 500 millilitre(s), closed with Screw cap (PolyPropylene)
(ID1) 250 millilitre(s): Box (Cardboard) with 1 Bottle (High Density PolyEthylene) with 250 millilitre(s), closed with Screw cap (PolyPropylene)

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
16/09/2014

Manufacturing sites for batch release:
Norbrook Laboratories (Ireland) Limited
Norbrook Laboratories Limited

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401978.00.00

Date of authorisation status change:
30/07/2019

Reference member state:
Germany

Procedure number:
DE/V/0320/001

Concerned member states:
Netherlands United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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