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VIRBAMEC Pour-On solution for cattle 5 mg/ml

Authorised

- Ivermectin

Product identification

Medicine name:
VIRBAMEC Pour-On solution for cattle 5 mg/ml

Active substance:
Ivermectin

Target species:
Cattle

Route of administration:
Pour-on use

Product details

Active substance and strength:
Ivermectin
0.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Pour-on solution

Withdrawal period by route of administration:

Pour-on use:

- **Cattle**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Size: 500 mlContainer: Opaque high density polyethylene bottle.Closure: Low density polyethylene seal fitted with a child resistant screw cap.Dosing device: separate measuring chamber, made of polyproylene, capable of delivering doses of 10 to 25 ml at 5 mlintervals.

Size: 1000 mlContainer: Opaque high density polyethylene bottle.Closure: Low density polyethylene seal fitted with a child resistant screw cap.Dosing device: separate measuring chamber, made of polyproylene, capable of delivering doses of 10 to 25 ml at 5 mlintervals

Size: 2.5 LContainer: Opaque high density polyethylene pack.Closure: Low density polyethylene seal fitted with a child resistant screw cap.Dosing device: made of polyproylene, to be used in conjunction with a dosing gun.

Size: 5.0 LContainer: Opaque high density polyethylene pack.Closure: Low density polyethylene seal fitted with a child resistant screw cap.Dosing device: made of polyproylene, to be used in conjunction with a dosing gun.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
15/05/2002

Manufacturing sites for batch release:
Virbac

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400568.00.00

Date of authorisation status change:
5/03/2009

Reference member state:
Ireland

Procedure number:
IE/V/0499/001

Concerned member states:
Belgium France Germany Luxembourg Portugal Spain
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents

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