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Chanectin 5 mg/ml pour-on solution

Authorised

- Ivermectin

Product identification

Medicine name:
Chanectin 5 mg/ml pour-on solution

Active substance:
Ivermectin

Target species:
Cattle

Route of administration:
Pour-on use

Product details

Active substance and strength:
Ivermectin
5.00 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

Package description:

High density polyethylene squeeze measure pour containers with child resistantclosures. Pack size: 500ml.
High density polyethylene squeeze measure pour containers with child resistantclosures. Pack size: 250ml.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure.The 1L pack will also have a dial a dose dosing cup.Pack size: 6L.The 6L consists of a 5L and 1L pack combined in one carton.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure.Pack size: 5L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure.Pack size: 2.5L.
High density polyethylene squeeze measure pour containers with child resistantclosures. Pack size: 1L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure.The 1L pack will also have a dial a dose dosing cup.Pack size: 1L.

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
14/11/2005

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400864.00.00

Date of authorisation status change:
9/09/2013

Reference member state:
Ireland

Procedure number:
IE/V/0170/001

Concerned member states:
Germany Portugal United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Combined File of all Documents