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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:

Ireland

Package description:

High density polyethylene squeeze measure pour containers with child resistantclosures. Pack size: 1L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure.Pack size: 2.5L.
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.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 squeeze measure pour containers with child resistantclosures. Pack size: 250ml.
High density polyethylene squeeze measure pour containers with child resistantclosures. Pack size: 500ml.
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:
3/12/2004

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/154/001

Date of authorisation status change:
3/12/2004

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

Combined File of all Documents