Anisec 0.5 % w/v Pour-on Solution

Authorised

- Ivermectin

Product identification

Medicine name:
Anisec 0.5 % w/v 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 container (flat bottomed flexi packs) with a 38mm tamperevident closure.Pack size: 1L. The 1L pack will also have a dial a dose dosing cup.
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.Pack sizes: 6L.The 6L consists of a 5L and 1L pack combined in one carton. The 1L pack will also have a dial a dose dosing cup.
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.

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
13/05/2005

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/158/001

Date of authorisation status change:
13/05/2005

Reference member state:
Ireland

Procedure number:
IE/V/0169/001

Concerned member states:
Belgium Germany Netherlands

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