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:
Belgium

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:
21/11/2005

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V277234

Date of authorisation status change:
21/11/2005

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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