

Anisec 0.5 % w/v Pour-on Solution

Authorised

- Ivermectin

Product identification

Medicine name:

Anisec 0.5 % w/v Pour-on Solution
Iverpour 0.5 % w/v Pour-on oplossing
Iverpour 0.5 % w/v Solution pour pour-on
Iverpour 0.5 % w/v Lösung zum übergießen

Active substance:

Ivermectin

Target species:

Cattle

Route of administration:

Topical 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:

Topical 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

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 resistant closures. Pack size: 500ml.

High density polyethylene squeeze measure pour containers with child resistant closures. 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 resistant closures. 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 Denmark Germany Netherlands

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

Documents

Summary of Product Characteristics

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Package Leaflet

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