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Animec 0.5% w/v Pour-on Solution

Authorised

- Ivermectin

Product identification

Medicine name:
Animec 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
0.50 gram(s) / 100.00 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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure (2.5L).Pack size: 2.5L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure (5L).Pack size: 5L.
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: 6L.The 6L consists of a 5L and 1L pack combined in one carton.
High density polyethylene squeeze measure pour containers with child resistant closures. Pack size: 250ml.
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: 1L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure (1L).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:
24/11/2005

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 08749/3140

Date of authorisation status change:
4/04/2022

Reference member state:
Ireland

Procedure number:
IE/V/0172/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents