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Sumex 5 mg/ml Pour on Solution for Cattle

Authorised

- Ivermectin

Product identification

Medicine name:
Sumex 5 mg/ml Pour on Solution for Cattle

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 resistant closures. Pack sizes: 1L

High density polyethylene squeeze measure pour containers with child resistantclosures. Pack sizes: 500ml

High density polyethylene squeeze measure pour containers with child resistant closures. Pack sizes: 250ml

High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure (1L and 5L).The 1L pack will also have a dial a dose dosing cup.Pack sizes: 6L.The 6L pack size consists of a 5L and 1L pack combined in one carton.

High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure (5L).Pack sizes: 5L.

High density polyethylene container (flat bottomed flexi packs) with a 38mm tamper evident closure (2.5L).Pack sizes: 2.5L

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 sizes: 1L

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
19/01/2007

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/160/001

Date of authorisation status change:
19/01/2007

Reference member state:
Ireland

Procedure number:
IE/V/0209/001

Concerned member states:
Greece Italy Spain

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