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Bovipour 5 mg/ml pour-on solution for cattle

Authorised

- Ivermectin

Product identification

Medicine name:
Bovipour 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 resistantclosures. Pack sizes: 1 L.
High density polyethylene squeeze measure pour containers with child resistantclosures. Pack sizes: 500 ml.
High density polyethylene squeeze measure pour containers with child resistantclosures. Pack sizes: 250 ml.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure.The 1 L pack will also have a dial a dose dosing cup.Pack sizes: 6 L.The 6L pack size consists of a 5 L and 1 L pack combined in one carton.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure (5 L).Pack sizes: 5 L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure (2.5 L).Pack sizes: 2.5 L.
High density polyethylene container (flat bottomed flexi packs) with a 38mm tamperevident closure (1 L).The 1 L pack will also have a dial a dose dosing cup.Pack sizes: 1 L.

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:
25/05/2007

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/162/001

Date of authorisation status change:
25/05/2007

Reference member state:
Ireland

Procedure number:
IE/V/0212/001

Concerned member states:
Austria 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