Animec 0.8mg/ml Oral Solution for Sheep

Authorised

- Ivermectin

Product identification

Medicine name:
Animec 0.8mg/ml Oral Solution for Sheep

Active substance:
Ivermectin

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Ivermectin
0.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Sheep**
  - Meat and offal. 10 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:

White flat-bottomed flexi packs (6L (5L + 1L)) composed of high density polyethylene container,with a 38 mm tamper evident polypropylene cap.Pack sizes (flexi pack): 6L (5L+1L)
Standard containers (jerri-cans) (1L) composed of high density polyethylene container, with tamperevident polyethylene cap.Pack sizes (jerri-cans): 1 Litre
White flat-bottomed flexi packs (2.5L) composed of high density polyethylene container,with a 38 mm tamper evident polypropylene cap.Pack sizes (flexi pack): 2.5 L
Standard containers (jerri-cans) (5 L) composed of high density polyethylene container, with tamperevident polyethylene cap.Pack sizes (jerri-cans): 5L
Standard containers (jerri-cans) (10L) composed of high density polyethylene container, with tamperevident polyethylene cap.Pack sizes (jerri-cans): 10 Litre
Standard containers (jerri-cans) (2.5L) composed of high density polyethylene container, with tamperevident polyethylene cap.Pack sizes (jerri-cans): 2.5 Litre
White flat-bottomed flexi packs (5L) composed of high density polyethylene container,with a 38 mm tamper evident polypropylene cap.Pack sizes (flexi pack): 5L.
White flat-bottomed flexi packs (1L) composed of high density polyethylene container,with a 38 mm tamper evident polypropylene cap.Pack size (flexi pack): 1 L

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
8/07/2010

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
VM 08749/4027

Date of authorisation status change:
19/11/2024

Reference member state:
Ireland

Procedure number:
IE/V/0249/001

Concerned member states:
Greece Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics