

Solantel 50mg/ml Oral Suspension for Sheep

Authorised

- Closantel sodium dihydrate

Product identification

Medicine name:

Solantel 50 mg/ml Oral Suspension for Sheep

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Active substance:

Closantel sodium dihydrate

Target species:

Sheep

Route of administration:

Oral use

Product details

Active substance and strength:

Closantel sodium dihydrate

54.38 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:**Oral use:**

-

Sheep

- Meat and offal. 56 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AG09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

Package description:

White high density polyethylene multidose container backpacks with high density polyethylene screw cap with induction-sealliners. Pack sizes:Box with 1 multidose container of 1 litre

White high density polyethylene multidose container backpacks with high density polyethylene screw cap with induction-sealliners. Pack sizes:Box with 1 multidose container of 2.5 litres

White high density polyethylene multidose container backpacks with high density polyethylene screw cap with induction-sealliners. Pack sizes:Box with 1 multidose container of 5 litres

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:

Norbrook Laboratories Limited

Marketing authorisation date:

4/08/2016

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 02000/4402

Date of authorisation status change:

23/01/2024

Reference member state:

Ireland

Procedure number:

IE/V/0552/001

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