

Tribex 5 % Oral Suspension for Sheep

Authorised

- Triclabendazole

Product identification

Medicine name:

Tribex 5 % Oral Suspension for Sheep

Active substance:

Triclabendazole

Target species:

Sheep

Route of administration:

Oral use

Product details

Active substance and strength:

Triclabendazole

50.00 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:

QP52AC01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Pack sizes: 1L pack contains 0.8L of product, Container: High density polyethylene Closure: Copolymer polypropylene with tamper evident seal Cap Liner: Polyfaced Steran Wad Spout: Polypropylene

Pack sizes: 5L pack contains 5L of product Container: High density polyethylene Closure: Copolymer polypropylene with tamper evident seal Cap Liner: Polyfaced Steran Wad Spout: Polypropylene

Pack sizes: 2.5L pack contains 2.2L of product Container: High density polyethylene Closure: Copolymer polypropylene with tamper evident seal Cap Liner: Polyfaced Steran Wad Spout: Polypropylene

Pack sizes: 2.5L pack contains 2.2L of product Container: High density polyethylene Closure: Copolymer polypropylene with tamper evident seal Cap Liner: Polyfaced Steran Wad Spout: Polypropylene

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

2/10/2000

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10987/146/001

Date of authorisation status change:

2/10/2000

Reference member state:

Ireland

Procedure number:

IE/V/0129/001

Concerned member states:

Netherlands 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