

Triclacert 10% Oral Suspension for Cattle

Authorised

- Triclabendazole

Product identification

Medicine name:

Triclacert 10% Oral Suspension for Cattle

Active substance:

Triclabendazole

Target species:

Cattle

Route of administration:

Oral use

Product details

Active substance and strength:

Triclabendazole

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

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

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Cattle

- Meat and offal. 56 day
- Milk. 84 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

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 2.5L 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:1L pack contains 1L of product Container: High density

polyethylene Closure: Copolymer polypropylene with tamper evident seal Cap Liner:

Polyfaced Steran Wad Spout: Polypropylene

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

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:

11/07/2007

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 08749/3046

Date of authorisation status change:

4/05/2024

Reference member state:

Ireland

Procedure number:

IE/V/0143/001

Concerned member states:

Czechia France Germany 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