Triclaben 100 mg/ml oral suspension for Cattle

Authorised

- Triclabendazole

Product identification

Medicine name:
Triclaben 100 mg/ml 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:

- **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

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Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Pack sizes:5L pack contains 5L of productContainer: High density polyethyleneClosure: Copolymer polypropylene with tamper evident sealCap Liner: Polyfaced Steran WadSpout: Polypropylene
Pack sizes:2.5L pack contains 2.5L of productContainer: High density polyethyleneClosure: Copolymer polypropylene with tamper evident sealCap Liner: Polyfaced Steran WadSpout: Polypropylene
Pack sizes:2.5L pack contains 2.2L of productContainer: High density polyethyleneClosure: Copolymer polypropylene with tamper evident sealCap Liner: Polyfaced Steran WadSpout: Polypropylene
Pack sizes:1L pack contains 1L of productContainer: High density polyethyleneClosure: Copolymer polypropylene with tamper evident sealCap Liner: Polyfaced Steran WadSpout: Polypropylene
Pack sizes:1L pack contains 0.8L of product Container: High density polyethyleneClosure: Copolymer polypropylene with tamper evident sealCap Liner: Polyfaced Steran WadSpout: Polypropylene

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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:
6/12/2002

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/059/001

Date of authorisation status change:
6/12/2002

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