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Curazole 50 mg/ml oral drench for cattle and sheep

Authorised

Fenbendazole

Product identification

Medicine name:

Curazole 50 mg/ml oral drench for cattle and sheep

Active substance:

Fenbendazole

Target species:

Cattle

Sheep

Route of administration:

Oral use

Product details

Active substance and strength:

Fenbendazole

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

Cattle

- Meat and offal. 28 day
- Milk. 120 hour

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Sheep

- Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

- 1 L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
- 5 L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
- 2.5 L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
- 2.5 L (back pack) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
- 1 L (flat bottom flexi) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.

Additional information

www.adrreports.eu/vet

Summary of Product Characteristics

Documents

Entitlement type: Marketing Authorisation
Legal basis of product authorisation: Informed consent (abridged application) - Council Directive 81/851/EEC
Marketing authorisation holder: Univet Limited
Marketing authorisation date: 12/03/1992
Manufacturing sites for batch release: Univet Limited
Responsible authority: Health Products Regulatory Authority
Authorisation number: VPA10990/015/003
Date of authorisation status change: 12/03/1992
To consult adverse reactions on veterinary medicinal products please go to