Ranide 30 mg/ml Oral Drench

Authorised

- Rafoxanide

Product identification

Medicine name:
Ranide 30 mg/ml Oral Drench

Active substance:
Rafoxanide

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Rafoxanide
30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**

- Meat and offal. 60 day

- **Sheep**

  - Meat and offal. 60 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AG05

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Ireland

Package description:
5L (flexipack), HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
2.5L (jerrican), HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
1L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
5L (jerrican), HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.
2.5L (backpack), HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner.

Additional information

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:
17/02/1995

Manufacturing sites for batch release:
Univet Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10990/027/001

Date of authorisation status change:
17/02/1995

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics
