Fenafluke 5% w/v Oral Suspension

Authorised

- Fenbendazole
- Rafoxanide

Product identification

Medicine name:
Fenafluke 5% w/v Oral Suspension

Active substance:
Fenbendazole
Rafoxanide

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Fenbendazole
50.00 milligram(s) / 1.00 millilitre(s)

Rafoxanide
50.00 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:
QP52AC13
QP52AX

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

Authorisation status:
Valid

Authorised in:
Ireland

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

1L (flexipack) HDPE white 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 application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharvet (Ireland) Limited

Marketing authorisation date:
18/09/1998

Manufacturing sites for batch release:
Pharvet (Ireland) Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10462/003/001

Date of authorisation status change:
18/09/1998

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

Documents

Summary of Product Characteristics