Curafluke 10% w/v Oral Drench

Authorised

- Fenbendazole
- Rafoxanide

Product identification

Medicine name:
Curafluke 10% w/v Oral Drench

Active substance:
Fenbendazole

Rafoxanide

Target species:
Cattle

Route of administration:
Oral use

Product details

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

Rafoxanide
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 60 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC30

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

5 L (backpack) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner. The product may be marketed with or without an outer carton.
2.5 L (back pack) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner. The product may be marketed with or without an outer carton.
1 L (back pack) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner. The product may be marketed with or without an outer carton.
5 L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner. The product may be marketed with or without an outer carton.
2.5 L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner. The product may be marketed with or without an outer carton.
1 L (jerrican) HDPE white rigid containers closed with a polypropylene screw cap with an induction heat seal liner. The product may be marketed with or without an outer

carton.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Univet Limited

Marketing authorisation date:
15/09/2000

Manufacturing sites for batch release:
Univet Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10990/032/002

Date of authorisation status change:
15/09/2000

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

Documents

Summary of Product Characteristics