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Zanil Fluke Drench

Not authorised

- Oxyclozanide

Product identification

Medicine name:
Zanil Fluke Drench

Active substance:
Oxyclozanide

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Oxyclozanide
3.40 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Milk. 72 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52A

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

Authorisation status:
Surrendered

Authorised in:
Ireland

Package description:
High density polyethylene back-packs of 5 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.
High density polyethylene back-packs of 2.5 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.
High density polyethylene back-packs of 1 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.
High density polyethylene bottles of 10 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.
High density polyethylene bottles of 5 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.
High density polyethylene bottles of 2.5 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.
High density polyethylene bottles of 1 litre nominal volume. Closure: Polypropylene or urea formaldehyde screw cap with wads of PVDC-faced paper on a pulpboard liner.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Intervet (Ireland) Limited

Marketing authorisation date:
1/10/1999

Manufacturing sites for batch release:
Trirx Segre

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/262/001

Date of authorisation status change:
14/05/2025

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

Documents

Summary of Product Characteristics