COCCIRIL 2.5 MG/ML ORAL SUSPENSION FOR CATTLE AND SHEEP

Authorised

- Diclazuril

Product identification

Medicine name:
COCCIRIL 2.5 MG/ML ORAL SUSPENSION FOR CATTLE AND SHEEP

Active substance:
Diclazuril

Target species:
Cattle (calf)
Sheep (lamb)

Route of administration:
Oral use

Product details

Active substance and strength:
Diclazuril
2.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle (calf)**
  - Meat and offal. 0 day

- **Sheep (lamb)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51BC03

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

Authorisation status:
Valid

Authorised in:
Finland

Package description:
White, high-density polyethylene back-pack bottle of 1 L closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 1 L back-pack bottle per cardboard box together with a spigot cap and back-pack strap.
White, high-density polyethylene back-pack bottle of 5 L closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 5 L back-pack bottle per cardboard box together with a spigot cap and back-pack strap.
White, high-density polyethylene hanging bottle of 250 mL closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 250 mL bottle per cardboard box together with a spigot cap.

White, high-density polyethylene back-pack bottle of 2,5 L closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 2.5 L back-pack bottle per cardboard box together with a spigot cap and back-pack strap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
19/03/2026

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Finnish Medicines Agency

Authorisation number:
43403

Date of authorisation status change:
19/03/2026

Reference member state:
France

Procedure number:
FR/V/0479/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland

Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

Generic of:
600000042080

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.