VECOXAN 2.5 MG/ML ORAL SUSPENSION

Authorised

- Diclazuril

Product identification

Medicine name:
VECOXAN 2.5 MG/ML ORAL SUSPENSION
Vecoxan 2,5 mg/ml suspensie voor oraal gebruik

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:

Netherlands

Package description:

1 container of 1l and accessory box containing spouted cap and harness
Cardboard box with 1 container of 200 ml and accessory box containing spouted cap and harness
1 container of 5l and accessory box containing spouted cap and harness
1 container of 2.5l and accessory box containing spouted cap and harness

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet Nederland B.V.

Marketing authorisation date:
8/02/2000

Manufacturing sites for batch release:
Intervet Productions S.A.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 9660

Date of authorisation status change:
26/04/2022

Reference member state:
France

Procedure number:
FR/V/0113/001

Concerned member states:
Austria Belgium Czechia Germany Greece Hungary Ireland Italy Luxembourg Netherlands Norway Portugal Slovakia Spain

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

Documents

Combined File of all Documents

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