VECOXAN 2.5 MG/ML ORAL SUSPENSION

Authorised

- Diclazuril

Product identification

Medicine name:
VECOXAN 2.5 MG/ML ORAL SUSPENSION

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

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Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Available only in French

Available only in French

Available only in French

Available only in French

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet

Marketing authorisation date:
20/07/1998

Manufacturing sites for batch release:
INTERVET PRODUCTIONS

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/5395150 6/1998

Date of authorisation status change:
20/07/2008

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

Summary of Product Characteristics

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

Package Leaflet and Labelling

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