

VECOXAN 2.5 MG/ML ORAL SUSPENSION FOR LAMBS AND CALVES

Authorised

- Diclazuril

Product identification

Medicine name:

VECOXAN 2.5 MG/ML ORAL SUSPENSION FOR LAMBS AND CALVES

VECOXAN, 2.5mg/ml, Perorální suspenze

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

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Sheep (lamb)

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51AJ03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

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 International B.V.

Marketing authorisation date:

17/04/2007

Manufacturing sites for batch release:

Intervet Productions S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/015/07-C

Date of authorisation status change:

7/04/2011

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

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Package Leaflet

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Labelling

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

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