Cardisure 3.5 mg/ml Oral Solution for Dogs

Authorised

- Pimobendan

Product identification

Medicine name:
Cardisure 3.5 mg/ml Oral Solution for Dogs
Cardisure 3,5 mg/ml Lösung zum Eingeben für Hunde

Active substance:
Pimobendan

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Pimobendan
3.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC01CE90

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Brown high-density polyethylene bottles fitted with white polypropylene child resistant caps, and low-density polyethylenesyringe adaptors.A low-density polyethylene oral dosing syringe with graduations is supplied with the product.Pack size:Carton box containing 1 bottle of 168 ml and a 3 ml dosing syringe
Brown high-density polyethylene bottles fitted with white polypropylene child resistant caps, and low-density polyethylenesyringe adaptors.A low-density polyethylene oral dosing syringe with graduations is supplied with the product.Pack size:Carton box containing 1 bottle of 42 ml and a 1.5 ml dosing syringe

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:

10/10/2019

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
839151

Date of authorisation status change:
10/10/2019

Reference member state:
Ireland

Procedure number:
IE/V/0421/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland Greece Hungary Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Sweden

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
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Labelling

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