File downloaded on 2025-12-18

**Source URL:** https://medicines.health.europa.eu/veterinary/en/60000036889

# Cardisure Flavoured 5 mg Tablets for Dogs

Authorised

Pimobendan

## Product identification

#### **Medicine name:**

Cardisure Flavoured 5 mg Tablets for Dogs Cardisure Flavoured 5 mg Comprimé

#### **Active substance:**

Pimobendan

## **Target species:**

Dog

#### Route of administration:

Oral use

# **Product details**

## **Active substance and strength:**

Pimobendan

5.00 milligram(s) / 1.00 Tablet

### **Pharmaceutical form:**

**Tablet** 

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

OC01CE90

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Luxembourg

## Package description:

20 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)

4 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)

25 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)

2 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)

10 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)

10 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)

50 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)

5 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)

# Additional information

# **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

# Marketing authorisation holder:

Dechra Regulatory B.V.

## Marketing authorisation date:

12/07/2011

# Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Genera d.d.

Responsible authority:
Ministry Of Health And Social Security
Authorisation number:
V/914/11/07/1060
Date of authorisation status change:
12/07/2011
Reference member state:
Netherlands
Procedure number:
NL/V/0280/003
Concerned member states:
Austria Belgium Denmark Finland France Germany Greece Ireland Italy Luxembourg Norway Poland Portugal Spain Sweden
United Kingdom (Northern Ireland)
To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet
Documents
Summary of Product Characteristics
Summary of Froduct Characteristics
This document does not exist in this language (English). You can find it in another
language below.
Combined File of all Documents

lf
----