File downloaded on 2026-11-24
**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000036928

# Cardisure Flavoured 1.25 mg Tablets for Dogs

Authorised

- Pimobendan

## Product identification

**Medicine name:**
Cardisure Flavoured 1.25 mg Tablets for Dogs

**Active substance:**
Pimobendan

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Pimobendan
1.25 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QC01CE90

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

**Authorisation status:**
Valid

**Authorised in:**
United Kingdom (Northern Ireland)

**Available in:**
United Kingdom (Northern Ireland)

**Package description:**
10 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
10 blisters per carton, 10 tablets per blister (Aluminium - Aluminium blister)
2 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
2 blisters per carton, 10 tablets per blister (Aluminium - Aluminium blister)
25 blisters per carton, 10 tablets per blister (Aluminium - Aluminium blister)
25 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
5 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
5 blisters per carton, 10 tablets per blister (Aluminium - Aluminium 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:**
Eurovet Animal Health B.V.

**Marketing authorisation date:**
9/08/2011

**Manufacturing sites for batch release:**
Dales Pharmaceuticals Limited
Eurovet Animal Health B.V.
Genera d.d.

**Responsible authority:**
The Veterinary Medicines Directorate

**Authorisation number:**
Vm 16849/3015

**Date of authorisation status change:**
25/03/2022

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0280/001

**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 |

Combined File of all Documents

107727 PAR.pdf