

# Cardisure Flavoured 1.25 mg Tablets for Dogs

Authorised

- Pimobendan

## Product identification

**Medicine name:**

Cardisure Flavoured 1.25 mg Tablets for Dogs

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**Active substance:**

Pimobendan

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Pimobendan  
1.25 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QC01CE90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Available in:**

Netherlands

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

12/07/2011

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**Manufacturing sites for batch release:**

Dales Pharmaceuticals Limited  
Eurovet Animal Health B.V.  
Genera d.d.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 107727

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**Date of authorisation status change:**

19/01/2022

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0280/001

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**Concerned member states:**

Austria Belgium Denmark Finland France Germany Greece Ireland Italy  
Luxembourg Norway Poland Portugal Spain Sweden  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 24/11/2025

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Summary of Product Characteristics

107727 PAR.pdf