

# Cardisure Flavoured 5 mg Tablets for Dogs

Authorised

- Pimobendan

## Product identification

**Medicine name:**

Cardisure Flavoured 5 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

5.00 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:**

Poland

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**Available in:**

Poland

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**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)

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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:**

Eurovet Animal Health B.V.

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**Marketing authorisation date:**

8/08/2011

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.  
Genera d.d.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2130

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**Date of authorisation status change:**

8/08/2011

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0280/003

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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

Summary of Product Characteristics

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

## Package Leaflet

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

## Combined File of all Documents

English (PDF)

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## Labelling

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