

# Cardisan 1.25 mg chewable tablets for dogs

Authorised

- Pimobendan

## Product identification

**Medicine name:**

Cardisan 1.25 mg chewable tablets for dogs

Cardisan 1.25 mg Comprimé à croquer

Cardisan 1.25 mg Kauwtablet

Cardisan 1.25 mg Kautablette

**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:**

Chewable tablet

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**Withdrawal period by route of administration:****Oral use:**

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**Dog**

- All relevant tissues. no withdrawal period      Not applicable

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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:**

Belgium

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**Package description:**

Cardbox with 12 Aluminium-OPA/Aluminium/PVC blisters containing 10 tablets each.  
Cardbox with 10 Aluminium-OPA/Aluminium/PVC blisters containing 10 tablets each.  
Cardbox with 9 Aluminium-OPA/Aluminium/PVC blisters containing 10 tablets each.  
Cardbox with 6 Aluminium-OPA/Aluminium/PVC blisters containing 10 tablets each.  
Cardbox with 3 Aluminium-OPA/Aluminium/PVC blisters containing 10 tablets each.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

3/02/2023

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V661127

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**Date of authorisation status change:**

3/02/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0380/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Malta Norway Poland Portugal Romania Slovakia Slovenia  
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

Labelling

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