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# Pimocard 2.5 mg Flavoured Tablets for Dogs

Authorised

- Pimobendan

## Product identification

**Medicine name:**

Pimocard 2.5 mg Flavoured 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  
2.50 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:**

Germany

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**Package description:**

5 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.  
2 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.  
5 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.  
25 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.  
25 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.  
2 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.  
10 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.  
10 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.

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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:**

27/11/2015

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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:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

402262.01.00

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**Date of authorisation status change:**

14/12/2017

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0283/002

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**Concerned member states:**

Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Hungary Ireland  
Italy Latvia Lithuania Portugal Romania Slovakia Slovenia Spain  
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.