# Pimocard 2.5 mg Flavoured Tablets for Dogs

Authorised

- Pimobendan

## Product identification

**Medicine name:**
Pimocard 2.5 mg Flavoured Tablets for Dogs

**Active substance:**
Pimobendan

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Pimobendan
2.50 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:**
Slovenia

**Available in:**
Slovenia

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

## 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:**
18/11/2015

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

**Responsible authority:**
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

**Authorisation number:**
MR/V/0516/002

**Date of authorisation status change:**
18/11/2015

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0283/002

**Concerned member states:**
Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Hungary Ireland Italy Latvia Lithuania Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to 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.

Labelling

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