# Vetmedin 10 mg Kautabletten für Hunde

Authorised

Pimobendan

## Product identification

#### **Medicine name:**

Vetmedin 10 mg Kautabletten für Hunde Pimobendan Boehringer Ingelheim Vetmedica 10 mg Tuggtablett

#### **Active substance:**

Pimobendan

### **Target species:**

Dog

#### Route of administration:

Oral use

## **Product details**

## **Active substance and strength:**

Pimobendan

10.00 milligram(s) / 1.00 Tablet

#### **Pharmaceutical form:**

Chewable tablet

## Withdrawal period by route of administration:

## Oral use: Dog Anatomical therapeutic chemical veterinary (ATCvet) codes: OC01CE90 Legal status of supply: Veterinary medicinal product subject to veterinary prescription **Authorisation status:** Valid Authorised in: Sweden Package description: Cardboard box containing 50 tablets in a polyethylene bottle, closed with a polypropylene child-resistant screw cap Additional information **Entitlement type:** Marketing Authorisation Legal basis of product authorisation: Hybrid application (Article 13(3) of Directive No 2001/82/EC) Marketing authorisation holder: Boehringer Ingelheim Vetmedica GmbH Marketing authorisation date: 7/11/2013 Manufacturing sites for batch release: Boehringer Ingelheim Vetmedica GmbH

Responsible authority:

**Swedish Medical Products Agency** 

Date of authorisation status change: 7/11/2013  Reference member state: Austria  Procedure number: AT/V/0006/004  Concerned member states: Belgium Bulgaria Cyprus Czechia Estonia France Greece Hungary Iceland Ireland Italy Latvia Liechtenstein Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)  To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet  Documents  Package Leaflet  This document does not exist in this language (English). You can find it in another language below.	Authorisation number: 48938
Austria  Procedure number:  AT/V/0006/004  Concerned member states: Belgium Bulgaria Cyprus Czechia Estonia France Greece Hungary Iceland Ireland Italy Latvia Liechtenstein Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)  To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet  Documents  Package Leaflet  This document does not exist in this language (English). You can find it in another language below.	_
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