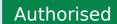
# Benazepril hydrochloride Le Vet 20 mg tablets for dogs



• Benazepril hydrochloride

## Product identification

#### **Medicine name:**

Benazepril hydrochloride Le Vet 20 mg tablets for dogs Benakor vet. 20 mg tabletter til hund

#### **Active substance:**

Benazepril hydrochloride

## **Target species:**

Dog

#### Route of administration:

Oral use

## **Product details**

## **Active substance and strength:**

Benazepril hydrochloride 20.00 milligram(s) / 1.00 Tablet

#### **Pharmaceutical form:**

**Tablet** 

## Withdrawal period by route of administration:

#### Oral use:

. Dog

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

OC09AA07

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### Authorised in:

Norway

#### Package description:

Carton box containing 5 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 6 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 5 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each Carton box containing 2 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each Carton box containing 1 PVC-PE-PVDC-Alu-foil blisters of 14 tablets Carton box containing 4 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each Carton box containing 3 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 6 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each Carton box containing 2 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 4 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 3 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each Carton box containing 7 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each Carton box containing 7 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 7 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 7 Alu-Alu -foil blisters blisters of 14 tablets each Carton box containing 1 Alu-Alu -foil blisters blisters of 14 tablets

## 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: Le Vet. B.V.
Marketing authorisation date: 2/03/2010
Manufacturing sites for batch release: Lelypharma B.V.
Responsible authority: Norwegian Medical Products Agency
Authorisation number: 07-4741
Date of authorisation status change: 30/01/2013
Reference member state: Netherlands
Procedure number: NL/V/0126/003
Concerned member states: Austria Belgium Czechia Estonia Finland France Ireland Italy Luxembourg Norway Poland Portugal Slovakia Spain
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents
Combined File of all 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.

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