# Benazepril hydrochloride Le Vet 20 mg tablets for dogs

Authorised

- Benazepril hydrochloride

## Product identification

**Medicine name:**
Benazepril hydrochloride Le Vet 20 mg tablets for dogs

**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

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QC09AA07

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Luxembourg

**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 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:**
Dechra Regulatory B.V.

**Marketing authorisation date:**
14/10/2009

**Manufacturing sites for batch release:**
Lelypharma B.V.

**Responsible authority:**
Ministry Of Health And Social Security

**Authorisation number:**
V 333/09/10/2033

**Date of authorisation status change:**
14/10/2009

**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 www.adrreports.eu/vet

## Documents

Combined File of all Documents

PuAR Benazepril Hydrochloride Le Vet - NL_V_0126_002-003_final clean.pdf