

# 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

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**Active substance:**

Benazepril hydrochloride

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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:**

Benazepril hydrochloride  
20.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QC09AA07

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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:**

Poland

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**Available in:**

Poland

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**Package description:**

Carton box containing 1 Alu-Alu-foil blisters blisters of 14 tablets

Carton box containing 7 Alu-Alu -foil blisters blisters of 14 tablets each

Carton box containing 7 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each

Carton box containing 3 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each

Carton box containing 4 Alu-Alu -foil blisters blisters of 14 tablets each

Carton box containing 2 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 3 Alu-Alu -foil blisters blisters of 14 tablets each

Carton box containing 4 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 2 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each

Carton box containing 5 PVC-PE-PVDC-Alu-foil blisters of 14 tablets each

Carton box containing 6 Alu-Alu -foil blisters blisters of 14 tablets each

Carton box containing 5 Alu-Alu -foil blisters blisters of 14 tablets each

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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:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

20/10/2008

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1838

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**Date of authorisation status change:**

20/10/2008

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0126/003

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**Concerned member states:**

Austria Belgium Czechia Estonia Finland France Ireland Italy Luxembourg  
Norway Poland Portugal Slovakia Spain

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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.

Labelling

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

Combined File of all Documents

PuAR Benazepril Hydrochloride Le Vet - NL\_V\_0126\_002-003\_final clean.pdf