

# 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:**

Luxembourg

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**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

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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:**

14/10/2009

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Ministry Of Health And Social Security

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**Authorisation number:**

V 333/09/10/2033

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**Date of authorisation status change:**

14/10/2009

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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

Combined File of all Documents

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