

Benazepril hydrochloride Le Vet 5 mg tablets for dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

Benazepril hydrochloride Le Vet 5 mg tablets for dogs

Active substance:

Benazepril hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Benazepril hydrochloride
5.00 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:

Netherlands

Package description:

Carton box containing 5 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 6 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 1 PVC-PE-PVDC-Alu-foil blisters of 14 tablets

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

Carton box containing 5 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 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 3 Alu-Alu -foil blisters blisters of 14 tablets each

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

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

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:

16/06/2008

Manufacturing sites for batch release:

Lelypharma B.V.

Genera d.d.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 10524

Date of authorisation status change:

19/01/2022

Reference member state:

Netherlands

Procedure number:

NL/V/0126/002

Concerned member states:

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

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www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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