Benazepril hydrochloride 5 mg tablets for dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Benazepril hydrochloride 5 mg tablets for dogs
Benazepril hydrochloride 5 mg tabletten voor honden

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 Piece

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**

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 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 3 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 2 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 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 3 Alu-Alu -foil blisters blisters of 14 tablets each
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 6 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 7 PVC-PE-PVDC-Alu-foil 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:
Le Vet. 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 10521

Date of authorisation status change:
19/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0125/002

Concerned member states:
Denmark Germany Greece Hungary Sweden

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents