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Benazepril hydrochloride 20 mg tablets for dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Benazepril hydrochloride 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:
Denmark

Package description:
Carton box containing 6 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 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 7 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 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 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 2 Alu-Alu -foil blisters blisters of 14 tablets each
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 3 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:
Dechra Regulatory B.V.

Marketing authorisation date:
4/04/2008

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
40766

Date of authorisation status change:
4/04/2008

Reference member state:
Netherlands

Procedure number:
NL/V/0125/003

Concerned member states:
Denmark Germany Greece Hungary Sweden

To consult adverse reactions on veterinary medicinal products please go to 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.

Combined File of all Documents

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