Cardalis 5 mg benazepril hydrochloride + 40 mg spironolactone - Chewable Tablet

Authorised

- Benazepril hydrochloride
- Spironolactone

Product identification

Medicine name:
Cardalis 5 mg benazepril hydrochloride + 40 mg spironolactone - Chewable Tablet

Active substance:
Benazepril hydrochloride

Spironolactone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Benazepril hydrochloride
5.00 milligram(s) / 1.00 Tablet

Spironolactone
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09BA07

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden

Package description:
Packaging:Bottle (HDPE), Package_size:90 tablets
Packaging:Bottle (HDPE), Package_size:30 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:
23/07/2012

Manufacturing sites for batch release:
Ceva Sante Animale
Catalent Germany Schorndorf GmbH
Ceva Sante Animale

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
23/07/2012

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

Documents

Combined File of all Documents

English (PDF)
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