

Cardalis 2.5 mg benazepril hydrochloride + 20 mg spironolactone - Chewable Tablet

Authorised

- Benazepril hydrochloride
- Spironolactone

Product identification

Medicine name:

Cardalis 2.5 mg benazepril hydrochloride + 20 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

2.50 milligram(s) / 1.00 Tablet

Spironolactone
20.00 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 , Latvia , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden

Package description:

Packaging:Bottle (HDPE), Package_size:30 tablets

Packaging:Bottle (HDPE), Package_size:90 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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