

BANACEP VET 5 MG FILM-COATED TABLETS FOR DOGS AND CATS

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

BANACEP VET 5 MG FILM-COATED TABLETS FOR DOGS AND CATS

Active substance:

Benazepril hydrochloride

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Benazepril hydrochloride

5.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated 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

Available in:

Netherlands

Package description:

Blister made of clear film of PVC/PE/PVDC and aluminium film containing 14 tablets.

Box with 1 blister (14 tablets).

Blister made of clear film of PVC/PE/PVDC and aluminium film containing 14 tablets.

Box with 10 blisters (140 tablets).

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:

Laboratorios Calier S.A.

Marketing authorisation date:

11/02/2008

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 10519

Date of authorisation status change:

22/03/2022

Reference member state:

France

Procedure number:

FR/V/0180/001

Concerned member states:

Belgium Germany Greece Ireland Italy Netherlands Poland Portugal
Romania Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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