Banacep Vet 20 mg film-coated tablets for dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Banacep Vet 20 mg film-coated 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:
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:
Ireland

Package description:
Box with 1 blister (14 tablets each)
Box with 10 blisters (14 tablets each)
Box with 4 blisters (14 tablets each)
Box with 2 blisters (14 tablets each)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Calier S.A.

Marketing authorisation date:
17/10/2025

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:

VPA10665/011/002

Date of authorisation status change:
17/10/2025

Reference member state:
France

Procedure number:
FR/V/0180/002

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

eu-puar-frv0180002-mr-rpe920-en.pdf