

Banacep Vet 20 mg Film-coated Tablet for Dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

Banacep Vet 20 mg Film-coated Tablet 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:

United Kingdom (Northern Ireland)

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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:

17/10/2011

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 20634/3005

Date of authorisation status change:

22/06/2024

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

13105_Banacep 20_PuAR 2025.pdf