

BANACEP VET 20 FILM-COATED TABLET FOR DOGS

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

BANACEP VET 20 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:

Greece

Available in:

Greece

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:

15/05/2012

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

2026/11-01-2017/K-0179602

Date of authorisation status change:

12/03/2019

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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

13105_Banacep 20_PuAR 2025.pdf