

SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT:

PRILBEN vet 5 film-coated tablet for dogs and cats [FR]
PRILBEN vet 5 mg film-coated tablet for dogs and cats [PL]
PRILBEN vet 5 mg film-coated tablet for dogs and cats
Benazepril hydrochloride [DE]
Prilben vet 5 mg film-coated tablet for dogs and cats
Benazepril hydrochloride [SE]
PRILBEN vet 5 mg film-coated tablet for dogs and cats
Benazepril hydrochloride [BE/DK/EL/ES/FI/IT/NL/PT/UK]

2. QUALITATIVE AND QUANTITATIVE COMPOSITION:

Each divisible tablet contains:

Benazepril 4.6 mg
(equivalent to Benazepril Hydrochloride 5 mg)
Excipients :
Titanium dioxide (E171)..... 1.929 mg
Iron oxide yellow (E172) 0.117 mg
Iron oxide red (E172)..... 0.014 mg
Iron oxide black (E172)..... 0.004 mg

For a full list of excipients, see section 6.1

3. PHARMACEUTICAL FORM:

Film-Coated Tablet
Beige oblong biconvex film-coated divisible tablets

4. CLINICAL PARTICULARS:

4.1 Target species:

Dogs and cats.

4.2 Indications for use, specifying target species:

Dogs:

Treatment of congestive heart failure.

Cats:

Reduction of proteinuria associated with chronic kidney disease.

4.3. Contraindications:

Do not use in case of hypersensitivity to the active substance or to any of the excipients.

Do not use in cases of hypotension, hypovolaemia, hyponatraemia or acute renal failure.

Do not use in cases of cardiac output failure due to aortic or pulmonary stenosis.
Do not use in pregnancy or lactation (see section 4.7).

4.4 Special warnings for each target species

None

4.5 Special precautions for use

Special precautions for use in animals

No evidence of renal toxicity to the veterinary medicinal product has been observed in dogs or cats during clinical trials, however, as is routine in cases of chronic kidney disease, it is recommended to monitor plasma creatinine, urea and erythrocyte counts during therapy.

The safety and the efficacy of the veterinary medicinal product has not been established in dogs and cats below 2.5 kg body weight.

Special precautions to be taken by the person administering the veterinary medicinal product to animals

Pregnant women should take special care to avoid accidental oral exposure, because angiotensin converting enzyme (ACE) inhibitors have been found to affect the unborn child during pregnancy in humans.

Wash hands after use.

In case of accidental oral ingestion, seek medical advice immediately and show the label or the package leaflet to the physician.

4.6 Adverse reactions (frequency and seriousness):

In double-blind clinical trials in dogs with heart failure, the product was well tolerated with an incidence of adverse reactions lower than observed in placebo-treated dogs.

A small number of dogs may exhibit transient vomiting, incoordination or signs of fatigue.

In cats and dogs with chronic kidney disease, the product may increase plasma creatinine concentrations at the start of therapy. A moderate increase in plasma creatinine concentrations following administrations of ACE inhibitors is compatible with the reduction in glomerular hypertension induced by these agents, and is therefore not necessarily a reason to stop therapy in the absence of other signs.

The product may increase food consumption and body weight in cats.

Emesis, anorexia, dehydration, lethargy and diarrhoea have been reported on rare occasions in cats.

4.7 Use during pregnancy and lactation

Do not use during pregnancy or lactation. The safety of the product has not been established in breeding, pregnant or lactating dogs and cats.

Benazepril reduced ovary / oviduct weights in cats when administered daily at 10 mg/kg body weight for 52 weeks.

Embryotoxic effects (foetal urinary tract malformations) were seen in trials with laboratory animals (rats) at maternally non-toxic doses.

4.8 Interaction with other medicaments and other forms of interaction:

In dogs with congestive heart failure, the product has been given in combination with digoxin, diuretics, pimobendan and anti-arrhythmic veterinary medicinal products without demonstrable adverse interactions.

. In humans, the combination of ACE inhibitors and Non-Steroidal Anti-Inflammatory Drugs (NSAIDs) can lead to reduced anti-hypertensive efficacy or impaired renal function. The combination of the product and other anti-hypertensive agents (e.g. calcium channel blockers, β -blockers or diuretics), anaesthetics or sedatives may lead to additive hypotensive effects. Therefore, concurrent use of NSAIDs or other medications with a hypotensive effect should be considered with care. Renal function and signs of hypotension (lethargy, weakness etc) should be monitored closely and treated as necessary.

Interactions with potassium preserving diuretics like spironolactone, triamterene or amiloride cannot be ruled out. It is recommended to monitor plasma potassium levels when using the product in combination with a potassium sparing diuretic because of the risk of hyperkalaemia.

4.9 Amounts to be administered and administration route:

Oral use

The product should be given orally once daily, with or without food. The duration of treatment is unlimited.

Dogs:

The product should be administered orally at a minimum dose of 0.25 mg (range 0.25-0.5) benazepril hydrochloride /kg bw once daily according to the following table

Weight of dog (kg)	Prilben Vet 5 mg Film-Coated Tablets	
	Standard dose	Double dose
> 5 - 10	0.5 tablet	1 tablet
>10 - 20	1 tablet	2 tablets

The dose may be doubled, still administered once daily, to a minimum of 0.5 mg/kg (range 0.5-1.0), if judged clinically necessary and advised by the veterinary surgeon.

Cats:

The product should be administered orally at a minimum dose of 0.5 mg (range 0.5-1.0) benazepril hydrochloride /kg bw once daily according to the following table :

Weight of cat (kg)	Prilben Vet 5 mg Film-Coated Tablets
2.5 – 5	0.5 tablet
> 5 - 10	1 tablet

4.10 Overdose (symptoms, emergency procedures, antidotes):

The product reduced erythrocyte counts in normal cats when dosed at 10 mg/kg body weight once daily for 12 months and in normal dogs when dosed at 150 mg/kg body weight once daily for 12 months, but this effect was not observed at the recommended dose during clinical trials in cats or dogs.

Transient reversible hypotension may occur in cases of accidental overdose. Therapy consists of intravenous infusion of warm isotonic saline.

4.11 Withdrawal period:

Not applicable

5 PHARMACOLOGICAL PROPERTIES

Pharmacotherapeutic group: ACE inhibitors, benazepril

ATCvet code: QC09AA07

5.1 Pharmacodynamic properties

Benazepril hydrochloride is a prodrug hydrolysed *in vivo* to its active metabolite, benazeprilat. Benazeprilat is a highly potent and selective inhibitor of ACE, thus preventing the conversion of inactive angiotensin I to active angiotensin II and thereby also reducing synthesis of aldosterone. Therefore, it blocks effects mediated by angiotensin II and aldosterone, including vasoconstriction of both arteries and veins, retention of sodium and water by the kidney and remodelling effects (including pathological cardiac hypertrophy and degenerative renal changes).

The product causes long-lasting inhibition of plasma ACE activity in dogs and cats, with more than 95% inhibition at peak effect and significant activity (>80% in dogs and >90% in cats) persisting 24 hours after dosing.

The product reduces the blood pressure and volume load on the heart in dogs with congestive heart failure.

In cats with experimental renal insufficiency, the product normalized the elevated glomerular capillary pressure and reduced the systemic blood pressure.

Reduction in glomerular hypertension may retard the progression of kidney disease by inhibition of further damage to the kidneys. Placebo controlled clinical field studies in cats with chronic kidney disease (CKD) have demonstrated that the product significantly reduced levels of urine protein and urine protein to creatinine ratio (UPC); this effect is probably mediated via reduced glomerular hypertension and beneficial effects on the glomerular basement membrane.

No effect of the product on survival in cats with CKD has been shown, but the product increased the appetite of the cats, particularly in more advanced cases.

5.2. Pharmacokinetic properties

After oral administration, of benazepril hydrochloride, peak levels of benazepril are attained rapidly (T_{max} 0.5 hour in dogs and within 2 hours in cats) and decline quickly as the active substance is partially metabolised by liver enzymes to benazeprilat. The systemic bioavailability is incomplete (~13% in dogs) due to incomplete absorption (38% in dogs, <30% in cats) and first pass metabolism. In dogs, peak benazeprilat concentrations (C_{max} of 37.6 ng/ml after a dose of 0.5 mg/kg benazepril hydrochloride) are achieved with a T_{max} of 1.25 hours. In cats, peak benazeprilat concentrations (C_{max} of 77.0ng/ml after a dose of 0.5 mg/kg benazepril hydrochloride) are achieved with a T_{max} of 2 hours.

Benazeprilat concentrations decline biphasically: the initial fast phase (with a $t_{1/2} = 1.7$ in dogs and $t_{1/2}$ of 2.4 hours in cats) represents elimination of free drug, while the terminal phase ($t_{1/2} = 19$ hours in dogs and $t_{1/2} = 29$ hours in cats) reflects the release of benazeprilat that was bound to ACE, mainly in the tissues.

Benazepril and benazeprilat are extensively bound to plasma proteins (85-90%), and in tissues are found mainly in the liver and kidney.

There is no significant difference in the pharmacokinetics of benazeprilat when benazepril hydrochloride is administered to fed or fasted dogs. Repeated administration of the veterinary medicinal product leads to slight bioaccumulation of benazeprilat ($R=1.47$ in dogs and $R=1.36$ in cats with 0.5 mg/kg), steady state being achieved within a few days (4 days in dogs).

Benazeprilat is excreted 54% via the biliary and 46% via the urinary route in dogs and 85% via the biliary and 15% via the urinary route in cats. The clearance of benazeprilat is not affected in dogs or cats with impaired renal function and therefore no adjustment of the product dose is required in either species in cases of renal insufficiency.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Iron oxide yellow (E-172)

Iron oxide red (E-172)

Iron oxide black (E-172)

Titanium dioxide (E-171)

Cellulose microcrystalline
Lactose monohydrate
Povidone
Maize starch
Silica colloidal anhydrous
Magnesium stearate
Hypromellose
Macrogol 8000

6.2 Incompatibilities

Not applicable

6.3 Shelf-life

Shelf life of the veterinary medicinal product as packaged for sale: 18 months
Return any halved tablet to the blister pack and use within 1 day. The blister pack should be inserted back into the cardboard box.

6.4 Special precautions for storage:

Do not store above 25°C. Store in a dry place.

6.5 Nature and composition of immediate packaging:

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

Box with:

- 1 blister (14 tablets)
- 10 blisters (140 tablets)

Not all pack size may be marketed.

6.6 Special precautions for the disposal of unused veterinary medicinal product or waste materials derived from the use of such products

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

7. MARKETING AUTHORISATION HOLDER

Chemo Ibérica, S.A.
Gran Vía Carlos III, 98 – 7^a
08028 Barcelona
SPAIN

8. MARKETING AUTHORISATION NUMBER

Vm 21690/4000

9. DATE OF THE FIRST AUTHORISATION

04/03/2008

10. DATE OF REVISION OF THE TEXT

December 2012

PROHIBITION OF SALE, SUPPLY AND/OR USE

Under veterinary prescription

Approved:  **21/12/2012**