Benamax Flavour 20 mg tablets for dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Benefortin Flavour 20 mg comprimidos para cães
Benamax Flavour 20 mg 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:
Tablet

Withdrawal period by route of administration:

Oral use:
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09AA07

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 1 blister strip of 7 tablets (7 tablets) Cardboard box with 2 blister strips of 7 tablets (14 tablets) Cardboard box with 4 blister strips of 7 tablets (28 tablets) Cardboard box with 10 blister strips of 7 tablets (70 tablets) PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 1 blister strip of 7 tablets (7 tablets) Cardboard box with 2 blister strips of 7 tablets (14 tablets) Cardboard box with 4 blister strips of 7 tablets (28 tablets) Cardboard box with 10 blister strips of 7 tablets (70 tablets) PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 1 blister strip of 7 tablets (7 tablets) Cardboard box with 2 blister strips of 7 tablets (14 tablets) Cardboard box with 4 blister strips of 7 tablets (28 tablets) Cardboard box with 10 blister strips of 7 tablets (70 tablets) PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 1 blister strip of 7 tablets (7 tablets) Cardboard box with 2 blister strips of 7 tablets (14 tablets) Cardboard box with 4 blister strips of 7 tablets (28 tablets) Cardboard box with 10 blister strips of 7 tablets (70 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:
Lavet Kft.

Marketing authorisation date:
28/10/2011

Manufacturing sites for batch release:
Lavet Pharmaceuticals Ltd.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
377/03/11RFVPT

Date of authorisation status change:
16/11/2022

Reference member state:
Hungary

Procedure number:
HU/V/0113/003

Concerned member states:
Austria Belgium Cyprus Czechia France Germany Ireland Latvia Lithuania Luxembourg Netherlands Poland Portugal Spain

United Kingdom (Northern Ireland)

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

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