Benamax Flavour 2.5 mg tablets for cats and dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Benamax Flavour 2.5 mg tablets for cats and dogs
Benefortin Flavour 2,5 mg Tabletten für Katzen und Hunde

Active substance:
Benazepril hydrochloride

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Benazepril hydrochloride
2.50 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC09AA07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Package description:

PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 14 tablets/blister. Cardboard box with 1 blister strip of 14 tablets (14 tablets) Cardboard box with 2 blister strips of 14 tablets (28 tablets) Cardboard box with 4 blister strips of 14 tablets (56 tablets) Cardboard box with 10 blister strips of 14 tablets (140 tablets)

PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 14 tablets/blister. Cardboard box with 1 blister strip of 14 tablets (14 tablets) Cardboard box with 2 blister strips of 14 tablets (28 tablets) Cardboard box with 4 blister strips of 14 tablets (56 tablets) Cardboard box with 10 blister strips of 14 tablets (140 tablets)

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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:
22/11/2011

Manufacturing sites for batch release:
Lavet Pharmaceuticals Ltd.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401601.00.00

Date of authorisation status change:
2/12/2016

Reference member state:
Hungary

Procedure number:
HU/V/0113/001

Concerned member states:
Austria Belgium Cyprus Czechia France Germany Greece 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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