Benamax Flavour 5 mg tablets for cats and dogs

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Benefortin Flavour 5 mg comprimés pour chats et chiens
Benamax Flavour 5 mg tablets for cats and dogs

Active substance:
Benazepril hydrochloride

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Benazepril hydrochloride
5.00 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

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

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

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:
19/01/2012

Manufacturing sites for batch release:
Lavet Kft.

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 993/12/01/2032

Date of authorisation status change:
19/01/2012

Reference member state:
Hungary

Procedure number:
HU/V/0113/002

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

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.
