# Benamax Flavour 20 mg tablets for dogs

• Benazepril hydrochloride

# **Product identification**

#### Medicine name:

Benamax Flavour 20 mg tablets for dogs Benefortin Flavour 20 mg tabletten voor honden

#### 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:

Netherlands

#### Package description:

PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. 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 4 blister strips of 7 tablets (28 tablets)
PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 2 blister strips of 7 tablets (14 tablets)
PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 2 blister strips of 7 tablets (14 tablets)
PVC/Aluminium/Polyamide blister -forming laminate with aluminium lidding foil with 7 tablets/blister. Cardboard box with 1 blister strips of 7 tablets (7 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:

#### 1/12/2011

#### Manufacturing sites for batch release:

Lavet Kft.

#### **Responsible authority:**

Medicines Evaluation Board

Authorisation number: REG NL 110136

#### Date of authorisation status change:

19/01/2022

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

## Documents

Combined File of all Documents

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