

Benazecare Flavour 5 mg Tablets for Dogs and Cats

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

Benazecare Flavour 5 mg Tablets for Dogs and Cats

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC09AA07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

Aluminium/aluminium blister packs containing 14 tablets, packed in a cardboard box with a package leaflet. BenazecareFlavour 5 mg tablets are supplied in packs of 14 tablets.

Aluminium/aluminium blister packs containing 14 tablets, packed in a cardboard box with a package leaflet. BenazecareFlavour 5 mg tablets are supplied in packs of 28 tablets.

Aluminium/aluminium blister packs containing 14 tablets, packed in a cardboard box with a package leaflet. BenazecareFlavour 5 mg tablets are supplied in packs of 56 tablets.

Aluminium/aluminium blister packs containing 14 tablets, packed in a cardboard box with a package leaflet. BenazecareFlavour 5 mg tablets are supplied in packs of 140 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:

Ecuphar

Marketing authorisation date:

22/10/2009

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V350716

Date of authorisation status change:

22/10/2009

Reference member state:

Ireland

Procedure number:

IE/V/0452/001

Concerned member states:

Austria Belgium France Germany Luxembourg Netherlands Spain

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 6/07/2025

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

Labelling

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