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Fortekor vet. 20 mg Tablett

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Fortekor vet. 20 mg Tablett

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09AA07

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Elanco GmbH

Marketing authorisation date:
16/12/2009

Manufacturing sites for batch release:
Novartis Sante Animale

Responsible authority:
Swedish Medical Products Agency

Authorisation number:

22907

Date of authorisation status change:
16/12/2009

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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