Benadil, 20mg, Film-coated tablet

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Benadil, 20mg, Film-coated tablet

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:
Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09AA07

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Aluminium/Aluminium Blister 7 x 14.0 Tablet
Aluminium/Aluminium Blister 2 x 14.0 Tablet
Aluminum/Plastic Blister 2 x 14.0 Tablet
Aluminum/Plastic Blister 7 x 14.0 Tablet

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:
Vetviva Richter GmbH

Marketing authorisation date:
15/11/2024

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:

72147

Date of authorisation status change:
15/11/2024

Reference member state:
Czechia

Procedure number:
CZ/V/0111/003

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Denmark France Germany Greece Hungary Luxembourg Netherlands Norway Poland Portugal Romania Slovakia 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

Labelling

Package Leaflet