Actikor 5 mg Film-coated Tablets for Dogs

Not authorised

- Benazepril hydrochloride

Product identification

Medicine name:
Actikor 5 mg Film-coated Tablets for Dogs
Actikor

Active substance:
Benazepril hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Benazepril hydrochloride
5.00 milligram(s) / 1.00 Piece

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:
Surrendered

Authorised in:
Sweden

Package description:
84 tablets in aluminium foil blister packs
56 tablets in aluminium foil blister packs
28 tablets in aluminium foil blister packs
140 tablets in aluminium foil blister packs
14 tablets in aluminium foil blister packs

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:
2/06/2010

Manufacturing sites for batch release:
Accord Healthcare Limited
Ecuphar

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
44445

Date of authorisation status change:
17/12/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0151/001

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