

Actikor 5 mg Film-coated Tablets for Dogs

Not
authorised

- Benazepril hydrochloride
- Benazepril hydrochloride

Product identification

Medicine name:

Actikor 5 mg Film-coated Tablets for Dogs

Active substance:

Benazepril hydrochloride

Benazepril hydrochloride

Target species:

Dog

Dog

Route of administration:

Oral use

Oral use

Product details

Active substance and strength:

Benazepril hydrochloride

5.00 milligram(s) / 1.00 Tablet

Benazepril hydrochloride
5.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated tablet

Withdrawal period by route of administration:

Oral use:

-

Dog

- Not applicable. no withdrawal period not applicable

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC09AA07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Netherlands

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:

Animalcare Limited

Marketing authorisation date:

7/06/2011

Manufacturing sites for batch release:

Accord Healthcare Limited

Ecuphar

Responsible authority:

Medicines Evaluation Board

Authorisation number:

107544

Date of authorisation status change:

16/10/2024

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