Vasotop P 1.25 mg tablets for dogs

Authorised

- Ramipril

Product identification

Medicine name:
Vasotop P 1.25 mg tablets for dogs
Vasotop P 1,25 mg tabletten voor honden

Active substance:
Ramipril

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Ramipril
1.25 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09AA05

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Carton box containing three 15 mL HD polyethylene containers containing 28 tablets closed by LD polypropylene tamper-evident child resistant screw cap. A desiccant capsule is inserted in the cap.
Carton box containing six 15 mL HD polyethylene containers containing 28 tablets closed by LD polypropylene tamper-evident child resistant screw cap. A desiccant capsule is inserted in the cap.
Carton box containing one 15 mL HD polyethylene container containing 28 tablets closed by LD polypropylene tamper-evident child resistant screw cap. A desiccant capsule is inserted in the cap.

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet Nederland B.V.

Marketing authorisation date:
17/06/2005

Manufacturing sites for batch release:
Intervet Ges.m.b.H.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 9608

Date of authorisation status change:
25/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0245/002

Concerned member states:
Austria Belgium Germany Greece Ireland Italy Luxembourg Norway Spain

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

Documents

Combined File of all Documents

English (PDF)
Published on: 17/02/2025
Download

PuAR Vastotop P 1.25mg - 24-6.pdf