Floxabactin 150 mg tablets for dogs

Authorised

- Enrofloxacin

Product identification

Medicine name:
Floxabactin 150 mg tablets for dogs

Active substance:
Enrofloxacin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Enrofloxacin
150.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Box with 2 blisters (20 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 2 blisters (20 tablets); Alu- PVC.PVDC blister with 10 tablets
Box with 15 blisters (150 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 15 blisters (150 tablets); Alu- PVC.PVDC blister with 10 tablets
Box with 10 blisters (100 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 10 blisters (100 tablets); Alu- PVC.PVDC blister with 10 tablets
Box with 1 blister (10 tablets); Alu-PVC.PVDC blister with 10 tablets
Box with 1 blister (10 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 3 blisters (30 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 5 blisters (50 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 3 blisters (30 tablets); Alu- PVC.PVDC blister with 10 tablets
Box with 5 blisters (50 tablets); Alu- PVC.PVDC blister with 10 tablets
Box with 6 blisters (60 tablets); Alu-PVC.PE.PVDC blister with 10 tablets
Box with 6 blisters (60 tablets); Alu- PVC.PVDC blister with 10 tablets

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:
Le Vet. B.V.

Marketing authorisation date:
7/07/2010

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
22/12/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0135/003

Concerned member states:
Austria Belgium Czechia Denmark Finland France Hungary Ireland Italy Poland Portugal Slovakia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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

Package Leaflet

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.

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