

Floxabactin 150 mg tablets for dogs

Authorised

- Enrofloxacin

Product identification

Medicine name:

Floxabactin 150 mg tablets for dogs

Floxabactin Vet. 150 mg tableter

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:

Denmark

Available in:

Denmark

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:

10/02/2011

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Danish Medicines Agency

Authorisation number:

44734

Date of authorisation status change:

10/02/2011

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.

Floxabactin PuAR NL-V-0135-001_003-DC final.pdf