

# Floxabactin 15 mg tablets for cats and dogs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Floxabactin 15 mg tablets for cats and dogs

Floxabactin Vet. 15 mg tableter

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**Active substance:**

Enrofloxacin

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**Target species:**

Dog

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Enrofloxacin

15.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01MA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Denmark

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**Available in:**

Denmark

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**Package description:**

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 5 blisters (50 tablets); Alu-PVC.PE.PVDC blister with 10 tablets

Box with 5 blisters (50 tablets); Alu- PVC.PVDC blister with 10 tablets

Box with 3 blisters (30 tablets); Alu-PVC.PE.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

Box with 3 blisters (30 tablets); Alu- PVC.PVDC blister with 10 tablets

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. B.V.

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**Marketing authorisation date:**

10/02/2011

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

44732

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**Date of authorisation status change:**

10/02/2011

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0135/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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