

# Enrotron Flavour 50 mg Tablets for dogs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Enrotron Flavour 50 mg Tablets for dogs  
ENROTRON FLAVOUR 50 mg tablete za pse

**Active substance:**

Enrofloxacin

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Enrofloxacin  
50.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Tablet

**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:**

Slovenia

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**Package description:**

(ID3) 30 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 30 Tablet

(ID9) 50 Tablet: unspecified outer container with 1 Blister (Aluminium; Polyvinylchlorid) with 50 Tablet

(ID6) 10 Tablet: unspecified outer container with 1 Blister (Aluminium; Polyvinylchlorid) with 10 Tablet

(ID2) 20 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 20 Tablet

(ID5) 100 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 100 Tablet

(ID7) 20 Tablet: unspecified outer container with 1 Blister (Aluminium; Polyvinylchlorid) with 20 Tablet

(ID8) 30 Tablet: unspecified outer container with 1 Blister (Aluminium; Polyvinylchlorid) with 30 Tablet

(ID4) 50 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 50 Tablet

(ID1) 10 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 10 Tablet

(ID10) 100 Tablet: unspecified outer container with 1 Blister (Aluminium; Polyvinylchlorid) with 100 Tablet

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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:**

aniMedica GmbH

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### **Marketing authorisation date:**

19/08/2010

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### **Manufacturing sites for batch release:**

aniMedica GmbH

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### **Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

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### **Authorisation number:**

DC/V/0110/001

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### **Date of authorisation status change:**

19/08/2010

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### **Reference member state:**

Germany

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### **Procedure number:**

DE/V/0137/001

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### **Concerned member states:**

Austria Belgium Denmark Finland Greece Hungary Iceland Ireland  
Luxembourg Netherlands Poland Slovenia 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

### 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.