

# Enrotron Flavour 150 mg Tablets for dogs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Enrotron Flavour 150 mg Tablets for dogs

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**Active substance:**

Enrofloxacin

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Enrofloxacin  
150.00 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:**

Poland

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**Package description:**

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

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

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

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

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

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

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

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

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

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

19/01/2011

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**Manufacturing sites for batch release:**

aniMedica GmbH

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2055

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**Date of authorisation status change:**

19/01/2011

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0137/002

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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  
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## Documents

### Summary of Product Characteristics

English (RTF)

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### Package Leaflet

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