

Enrotron Flavour 150 mg Tablets for dogs

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enrotron Flavour 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:

Poland

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

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:

aniMedica GmbH

Marketing authorisation date:

19/01/2011

Manufacturing sites for batch release:

aniMedica GmbH

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2055

Date of authorisation status change:

19/01/2011

Reference member state:

Germany

Procedure number:

DE/V/0137/002

Concerned member states:

Austria Belgium Denmark Finland Greece Hungary Iceland Ireland
Luxembourg Netherlands Poland Slovenia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (RTF)

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Package Leaflet

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