Enrotron Flavour 150 mg Tablets for dogs

Not 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:
Surrendered

Authorised in:
Belgium

Package description:
(ID10) 100 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 100 Tablet
(ID9) 50 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 50 Tablet
(ID8) 30 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 30 Tablet
(ID7) 20 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 20 Tablet
(ID6) 10 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) 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:
2/04/2012

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
20/11/2024

Reference member state:
Germany

Procedure number:
DE/V/0137/002

To consult adverse reactions on veterinary medicinal products please go to 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.