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

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Slovenia

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

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:
19/08/2010

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0110/001

Date of authorisation status change:
19/08/2010

Reference member state:
Germany

Procedure number:
DE/V/0137/001

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

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.