

Enrobactin 25 mg/ml concentrate for oral solution for pet rabbits, rodents, ornamental birds and reptiles

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enrobactin 25 mg/ml concentrate for oral solution for pet rabbits, rodents, ornamental birds and reptiles

Active substance:

Enrofloxacin

Target species:

Reptile

Ornamental bird

Rabbit

Rodents

Route of administration:

Oral use

Product details

Active substance and strength:

Enrofloxacin
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Concentrate for oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:

30 ml amber type III glass bottles closed with tamper-evident HDPE/LDPE screw caps with ring and colourless LDPE syringe (5 ml dosing syringe) insert in a carton box containing 30 ml

10 x (1x50 ml) amber type III glass bottles closed with tamper-evident HDPE/LDPE screw caps with ring and colourless LDPE syringe (5 ml dosing syringe) insert in a carton box containing 50 ml

50 ml amber type III glass bottles closed with tamper-evident HDPE/LDPE screw caps with ring and colourless LDPE syringe (5 ml dosing syringe) insert in a carton box containing 50 ml

10 x 1x30 ml amber type III glass bottles closed with tamper-evident HDPE/LDPE screw caps with ring and colourless LDPE syringe (5 ml dosing syringe) insert in a carton box containing 30 ml

10 x 1x10 ml amber type III glass bottles closed with tamper-evident HDPE/LDPE screw caps with ring and colourless LDPE syringe (1 ml dosing syringe) insert in a carton box containing 10 ml

10 ml amber type III glass bottles closed with tamper-evident HDPE/LDPE screw caps with ring and colourless LDPE syringe (1 ml dosing syringe) insert in a carton box containing 10 ml

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:

Le Vet. Beheer B.V.

Marketing authorisation date:

5/04/2017

Manufacturing sites for batch release:

Floris Veterinaire Producten B.V.

Responsible authority:

Finnish Medicines Agency

Authorisation number:

32541

Date of authorisation status change:

5/04/2017

Reference member state:

Netherlands

Procedure number:

NL/V/0315/001

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Denmark Finland France Germany
Greece Hungary Ireland Italy Luxembourg Norway Poland Portugal
Romania Slovakia Slovenia Sweden United Kingdom (Northern Ireland)

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