Orniflox 25 mg/ml Oral Solution for small mammals, birds and reptiles

Authorised

- Enrofloxacin

Product identification

Medicine name:
Orniflox 25 mg/ml Oral Solution for small mammals, birds and reptiles
Orniflox 25 mg/ml Concentraat voor drank
Orniflox 25 mg/ml Solution à diluer pour solution buvable
Orniflox 25 mg/ml Konzentrat zur Herstellung einer Lösung zum Einnehmen

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

Withdrawal period by route of administration:
Oral use:

- **Reptile**
- **Ornamental bird**
- **Rabbit**
- **Rodents**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
1 x 10 ml Type III amber glass bottles of 10 ml closed with tamper-evident HDPE screw caps with ring and colourless 1 ml dosing LDPE syringe insert in a carton box.
1 x 50 ml Type III amber glass bottles of 50 ml closed with tamper-evident HDPE screw caps with ring and colourless 5 ml dosing LDPE syringe insert in a carton box.
10 x (1 x 50 ml) Type III amber glass bottles of 50 ml closed with tamper-evident HDPE screw caps with ring and colourless 5 ml dosing LDPE syringe insert in a carton

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

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:
Avimedical B.V.

Marketing authorisation date:
23/11/2015

Manufacturing sites for batch release:
Floris Veterinaire Produkten B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V483057

Date of authorisation status change:
23/11/2015

Reference member state:
Netherlands

Procedure number:
NL/V/0294/001

Concerned member states:
Belgium France Germany Ireland Luxembourg

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.
