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 SOLUTION A DILUER POUR SOLUTION BUVABLE POUR LAPINS DE COMPAGNIE, RONGEURS, OISEAUX D'ORNEMENT ET 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

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:
France

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:
9/09/2015

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1078206 0/2015

Date of authorisation status change:
17/12/2020

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.
