

# Orniflox 25 mg/ml Concentrate for Oral Solution for Pet Rabbits, Rodents, Ornamental Birds and Reptiles

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Orniflox 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:**

United Kingdom (Northern Ireland)

---

**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:**

2/12/2015

---

**Manufacturing sites for batch release:**

Floris Veterinaire Producten B.V.

---

**Responsible authority:**

The Veterinary Medicines Directorate

---

**Authorisation number:**

Vm 43564/4001

---

**Date of authorisation status change:**

1/07/2024

---

**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](http://www.adrreports.eu/vet)