

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

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:

Germany

Available in:

Germany

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:

3/09/2015

Manufacturing sites for batch release:

Floris Veterinaire Producten B.V.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402150.00.00

Date of authorisation status change:

12/02/2021

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

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.