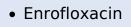
Orniflox 25 mg/ml Oral Solution for small mammals, birds and reptiles



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

Authorised

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

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

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