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# Quinoflox 100 mg/ml solution for use in drinking water, chicken and rabbits

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**
Quinoflox 100 mg/ml solution for use in drinking water, chicken and rabbits

**Active substance:**
Enrofloxacin

**Target species:**
Rabbit
Chicken

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Enrofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Oral solution

**Withdrawal period by route of administration:**

**Oral use:**

- **Rabbit**
  - Meat and offal. 2 day

- **Chicken**
  - Meat and offal. 4 day

Eggs: Do not use in birds producing eggs for human consumption

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01MA90

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

**Authorisation status:**
Valid

**Authorised in:**
Portugal

**Available in:**
Portugal

**Package description:**
1 litre white high density polyethylene containers, provided with green screw seal cap of the same material, with an aluminium disk sealed by induction.
4 X 5 litre white high density polyethylene containers, provided with green screw seal cap of the same material, with an aluminium disk sealed by induction
5 litre white high density polyethylene containers, provided with green screw seal cap of the same material, with an aluminium disk sealed by induction
12 X 1 litre white high density polyethylene containers, provided with green screw seal cap of the same material, with an aluminium disk sealed by induction

## 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:**
Global Vet Health S.L.

**Marketing authorisation date:**
8/08/2011

**Manufacturing sites for batch release:**
S P Veterinaria S.A.

**Responsible authority:**
Directorate General For Food And Veterinary

**Authorisation number:**
367/001/11DFVPT

**Date of authorisation status change:**
20/04/2022

**Reference member state:**
Portugal

**Procedure number:**
PT/V/0144/001

**Concerned member states:**
Belgium France Germany Italy Poland Romania
United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents