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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:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

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:
15/08/2011

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 36167/4001

Date of authorisation status change:
31/01/2023

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