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
NYOFLOX 100 MG/ML SOLUTION POUR ADMINISTRATION DANS L'EAU DE BOISSON POULETS ET LAPINS

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:
France

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:
15/12/2011

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7845975 5/2011

Date of authorisation status change:
1/08/2016

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

Summary of Product Characteristics

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

Package Leaflet and Labelling

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

Combined File of all Documents