File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000091716

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:
Belgium

Available in:
Belgium

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:
28/09/2011

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V400434

Date of authorisation status change:
28/09/2011

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

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.

Combined File of all Documents

Labelling

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