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# Quinoflox 100 mg/ml solution for injection for cattle and pigs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**
Quinoflox 100 mg/ml solution for injection for cattle and pigs

**Active substance:**
Enrofloxacin

**Target species:**
Pig
Cattle

**Route of administration:**
Intramuscular use
Intravenous use
Subcutaneous use

## Product details

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

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**

**Intramuscular use:**

- **Pig**
  - Meat and offal. 13 day

**Intravenous use:**

- **Cattle**
  - Meat and offal. 5 day
  - Milk. 72 hour

**Subcutaneous use:**

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 96 hour

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

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

**Authorisation status:**
Valid

**Authorised in:**
Romania

**Available in:**
Romania

**Package description:**

Amber polypropylene vial of 50 ml provided with a grey rubber-butyl stopper and aluminium seal with a green Flip-Off sealing.
Amber polypropylene vial of 100 ml provided with a grey rubber-butyl stopper and aluminium seal with a green Flip-Off sealing.
Amber polypropylene vial of 250 ml provided with a pink (250 ml) rubber-butyl stopper and aluminium seal with a green Flip-Off sealing.

## 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:**
22/11/2012

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

**Responsible authority:**
Institute For Control Of Biological Products And Veterinary Medicines

**Authorisation number:**
170263

**Date of authorisation status change:**
6/03/2023

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0290/001

**Concerned member states:**
Bulgaria Croatia Poland Portugal Romania United Kingdom (Northern Ireland)

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

## Documents

Combined File of all Documents

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

Summary of Product Characteristics