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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:
Poland

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:
4/06/2014

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2366

Date of authorisation status change:
4/06/2014

Reference member state:
Ireland

Procedure number:
IE/V/0290/001

Concerned member states:
Bulgaria Croatia Poland Portugal 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

English (PDF)
Published on: 16/01/2026
Updated on: 14/03/2026
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Labelling

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Package Leaflet

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