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QUINOLCEN 100 mg/ml Injection

Authorised

- Enrofloxacin

Product identification

Medicine name:
QUINOLCEN 100 mg/ml Injection

Active substance:
Enrofloxacin

Target species:
Cattle
Sheep
Goat
Pig

Route of administration:
Subcutaneous use
Intravenous use
Intramuscular 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:

Subcutaneous use:

- **Cattle**
  - Meat and offal. no withdrawal period

Meat and offal: IV: 5 days. Via SC: 12 days

- **Sheep**
  - Meat and offal. 4 day

- **Goat**
  - Meat and offal. 6 day

- **Cattle**
  - Milk. no withdrawal period

Milk: Via IV: 3 days. Via SC: 4 days;

- **Sheep**
  - Milk. 3 day

- **Goat**
  - Milk. 4 day

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period

Meat and offal: IV: 5 days. Via SC: 12 days

-

Cattle

- Milk. no withdrawal period — Milk: Via IV: 3 days. Via SC: 4 days;

Intramuscular use:

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Pig

- Meat and offal. 13 day

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:
box containing 1 vial of 250 ml
box containing 1 vial of 100 ml
box containing 1 vial of 50 ml

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:

S P Veterinaria S.A.

Marketing authorisation date:
11/05/2010

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

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

Authorisation number:
150242

Date of authorisation status change:
6/03/2023

Reference member state:
Spain

Procedure number:
ES/V/0150/001

Concerned member states:
Belgium Bulgaria Greece Hungary Italy Luxembourg Poland Portugal Romania

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 14/03/2026

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Package Leaflet

Combined File of all Documents