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Unisol 100 mg/ml solution for injection for cattle and pigs

Not authorised

- Enrofloxacin

Product identification

Medicine name:
Unisol 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. 3 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Authorisation status:
Surrendered

Authorised in:
Austria

Package description:
Type II Amber glass vial of 100 ml capacity closed with grey bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 1 vial of 100 ml

Type II Amber glass vial of 100 ml capacity closed with grey bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 12 vials of 100 ml
Type II Amber glass vial of 250 ml capacity closed with pink bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 1 vial of 250 ml
Type II Amber glass vial of 250 ml capacity closed with pink bromobutyl rubber stopper and aluminium flip-off seal.Cardboard box containing 12 vials of 250 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:
Vetpharma Animal Health S.L.

Marketing authorisation date:
6/07/2011

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
Chemical Iberica Productos Veterinarios S.L.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00973

Date of authorisation status change:
11/12/2025

Reference member state:
Ireland

Procedure number:
IE/V/0255/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 29/09/2024
Updated on: 14/03/2026
Download

Package Leaflet

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Labelling

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