# ENROVET 100 mg/ml SOLUCION INYECTABLE

Not authorised

- Enrofloxacin

## Product identification

**Medicine name:**
ENROVET 100 mg/ml SOLUCION INYECTABLE

**Active substance:**
Enrofloxacin

**Target species:**
Cattle
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

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Cattle**
  - Meat and offal. no withdrawal period: Carne: Vía SC 12 días. Vía IV 5 días
  - Milk. no withdrawal period: Leche: Vía SC 4 días; Vía IV 3 días
- **Cattle**
  - Meat and offal. no withdrawal period: Carne: Vía SC 12 días. Vía IV 5 días
  - Milk. no withdrawal period: Leche: Vía SC 4 días; Vía IV 3 días

**Intravenous use:**

- **Cattle**
  - Meat and offal. no withdrawal period: Carne: Vía SC 12 días. Vía IV 5 días
  - Milk. no withdrawal period: Leche: Vía SC 4 días; Vía IV 3 días
- **Cattle**
  - Meat and offal. no withdrawal period: Carne: Vía SC 12 días. Vía IV 5 días
  - Milk. no withdrawal period: Leche: Vía SC 4 días; Vía IV 3 días

**Intramuscular use:**

- **Pig**
  - Meat and offal. 3 day

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

QJ01MA90

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

**Authorisation status:**
Surrendered

**Authorised in:**
Spain

**Package description:**
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Informed consent application (Article 13c of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Divasa Farmavic S.A.

**Marketing authorisation date:**
26/09/2000

**Manufacturing sites for batch release:**
Laboratorios Syva S.A.

**Responsible authority:**
Spanish Agency Of Medicines And Medical Devices

**Authorisation number:**
1339 ESP

**Date of authorisation status change:**
29/01/2026

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

## Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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