

# ENROSYVA 100 mg/ml

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

ENROSYVA 100 mg/ml

ENROSYVA 100MG/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ

**Active substance:**

Enrofloxacin

**Target species:**

Cattle

Pig

**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:****• Cattle**

- Meat and offal. no withdrawal period

Meat and offal: 12 days (sc) / 5 days (iv)

- Milk. no withdrawal period  
Milk: 4 days (sc) / 3 days (iv)

**• Pig**

- Meat and offal. 13 day

**Intravenous use:****• Cattle**

- Meat and offal. no withdrawal period

Meat and offal: 12 days (sc) / 5 days (iv)

- Milk. no withdrawal period  
Milk: 4 days (sc) / 3 days (iv)

**• Pig**

- Meat and offal. 13 day

**Subcutaneous use:****• Cattle**

- Meat and offal. no withdrawal period

Meat and offal: 12 days (sc) / 5 days (iv)

- Milk. no withdrawal period  
Milk: 4 days (sc) / 3 days (iv)

**• Pig**

- Meat and offal. 13 day

---

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

QJ01MA90

---

**Legal status of supply:**

This information is not available for this product.

---

**Authorisation status:**

Valid

---

**Authorised in:**

Greece

---

**Available in:**

Greece

---

**Package description:**

polypropylene vial of 250 ml

box containing 1 glass vial of 100 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:**

Laboratorios Syva S.A.

---

**Marketing authorisation date:**

10/07/2018

---

**Manufacturing sites for batch release:**

Laboratorios Syva S.A.U.

---

**Responsible authority:**

National Organization For Medicines

---

**Authorisation number:**

105063/08-11-2021/K-0233201

---

**Date of authorisation status change:**

7/11/2021

---

**Reference member state:**

Spain

---

**Procedure number:**

ES/V/0247/001

---

**Concerned member states:**

Greece Italy Poland Portugal

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Package Leaflet

Labelling

eu-PUAR-enrosyva-100-mg-ml-en.pdf

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000038215>