

# ENROSYVA 100 mg/ml solution for injection for cattle and pigs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

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

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**Active substance:**

Enrofloxacin

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**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)

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**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)

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**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)

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## **Pig**

- Meat and offal. 13 day

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

QJ01MA90

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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### **Authorised in:**

Spain

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### **Package description:**

box containing 1 glass vial of 100 ml

box containing 1 polypropylene vial of 250 ml

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## **Additional information**

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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### **Marketing authorisation holder:**

Laboratorios Syva S.A.

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### **Marketing authorisation date:**

18/10/2016

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### **Manufacturing sites for batch release:**

Laboratorios Syva S.A.

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### **Responsible authority:**

**Authorisation number:**

3487 ESP

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**Date of authorisation status change:**

18/10/2016

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0247/001

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**Concerned member states:**

Greece Italy Poland Portugal

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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

English (PDF)

Published on: 11/04/2023

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### Package Leaflet

English (PDF)

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Labelling

English (PDF)

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Combined File of all Documents

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