

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

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:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

box containing 1 glass vial of 100 ml

box containing 1 polypropylene vial 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:

Laboratorios Syva S.A.

Marketing authorisation date:

2/09/2016

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1049/01/16DFVPT

Date of authorisation status change:

24/04/2026

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

Documents

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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Summary of Product Characteristics

Package Leaflet

Labelling

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