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:
Greece

Available in:
Greece

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:
10/07/2018

Manufacturing sites for batch release:

Laboratorios Syva S.A.

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 11/04/2023
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Package Leaflet

English (PDF)

Published on: 11/04/2023

Download

Labelling

Combined File of all Documents

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