

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

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

CENFLOX 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:**

Subcutaneous use

Intravenous use

Intramuscular 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:**

**Subcutaneous use:**

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: s.c.: 14 days/i.v.: 7 days

- Milk. no withdrawal period Milk: sc: 120 hours (5 days)/i.v 72 hours (3 days)

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: s.c.: 14 days/i.v.: 7 days

- Milk. no withdrawal period Milk: sc: 120 hours (5 days)/i.v 72 hours (3 days)

**Intravenous use:**

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: s.c.: 14 days/i.v.: 7 days

- Milk. no withdrawal period Milk: sc: 120 hours (5 days)/i.v 72 hours (3 days)

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: s.c.: 14 days/i.v.: 7 days

- Milk. no withdrawal period Milk: sc: 120 hours (5 days)/i.v 72 hours (3 days)

**Intramuscular use:**

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**Pig**

- Meat and offal. 12 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:**

Romania

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**Package description:**

box containing 1 glass vial of 100 ml  
box containing 1 glass vial of 250 ml  
box containing 10 glass vials of 100 ml  
box containing 10 glass vials of 250 ml  
box containing 1 polypropylene vial of 100 ml  
box containing 1 polypropylene vial of 250 ml  
box containing 10 polypropylene vials of 100 ml  
box containing 10 polypropylene vials 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:**

Cenavisa S.L.

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**Marketing authorisation date:**

21/05/2019

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**Manufacturing sites for batch release:**

Cenavisa S.L.

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

240058

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**Date of authorisation status change:**

14/04/2025

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0296/001

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**Concerned member states:**

Bulgaria Croatia Czechia Hungary Latvia Lithuania Portugal Romania

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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

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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