

# 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:**

Spain

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**Package description:**

Available only in [Spanish](#)

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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:**

13/03/2019

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**Manufacturing sites for batch release:**

Cenavisa S.L.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3749 ESP

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**Date of authorisation status change:**

13/03/2019

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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

English (PDF)

Published on: 12/04/2023

[Download](#)

Combined File of all Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

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