

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

Active substance:

Enrofloxacin

Target species:

Cattle

Pig

Route of administration:

Subcutaneous use

Intravenous use

Intramuscular 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:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

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

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:

Cenavisa S.L.

Marketing authorisation date:

21/05/2019

Manufacturing sites for batch release:

Cenavisa S.L.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

240058

Date of authorisation status change:

14/04/2025

Reference member state:

Spain

Procedure number:

ES/V/0296/001

Concerned member states:

Bulgaria Croatia Czechia Hungary Latvia Lithuania Portugal Romania

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)

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