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:

- **Cattle**
  - Meat and offal. no withdrawal period

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

- **Cattle**
  - Milk. no withdrawal period

Milk: sc: 120 houras (5 days)/i.v 72 hours (3 days)

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period

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

- **Cattle**
  - Milk. no withdrawal period

Milk: sc: 120 houras (5 days)/i.v 72 hours (3 days)

Intramuscular use:

- **Pig**
  - Meat and offal. 12 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Portugal

Package description:
box containing 10 polypropylene vials of 250 ml
box containing 10 polypropylene vials of 100 ml
box containing 1 polypropylene vial of 250 ml
box containing 1 polypropylene vial of 100 ml
box containing 10 glass vials of 250 ml
box containing 10 glass vials of 100 ml
box containing 1 glass vial of 250 ml
box containing 1 glass vial of 100 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:
18/02/2019

Manufacturing sites for batch release:
Cenavisa S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1242/01/19DFVPT

Date of authorisation status change:
22/04/2025

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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