UNOFLOX 100 mg/ml solution for injection.

Not authorised

- Enrofloxacin

Product identification

Medicine name:
UNOFLOX 100 mg/ml solution for injection.

Active substance:
Enrofloxacin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous 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: s.c.: 14 days/i.v.: 7 days

  - Milk. no withdrawal period

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

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

Subcutaneous use:

- **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 houras (5 days)/i.v 72 hours (3 days)

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

Intravenous use:

- **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 houras (5 days)/i.v 72 hours (3 days)

- **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:
Revoked

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 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:
S P Veterinaria S.A.

Marketing authorisation date:

25/03/2019

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1249/01/19DFVPT

Date of authorisation status change:
1/01/2023

Reference member state:
Spain

Procedure number:
ES/V/0299/001

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