

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:**

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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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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 hours (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 hours (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