

Quinoflox 100 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Enrofloxacin

Product identification

Medicine name:

Quinoflox 100 mg/ml solution for injection for cattle and pigs

Quinoflox 100 mg/ml Solution for Injection for Cattle and Pigs

Active substance:

Enrofloxacin

Target species:

Pig

Cattle

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous 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:**

-

Pig

- Meat and offal. 13 day

Intravenous use:

-

Cattle

- Meat and offal. 5 day

- Milk. 72 hour

Subcutaneous use:

-

Cattle

- Meat and offal. 12 day

- Milk. 96 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

Amber polypropylene vial of 50 ml provided with a grey rubber-butyl stopper and aluminium seal with a green Flip-Off sealing.

Amber polypropylene vial of 100 ml provided with a grey rubber-butyl stopper and aluminium seal with a green Flip-Off sealing.

Amber polypropylene vial of 250 ml provided with a pink (250 ml) rubber-butyl stopper and aluminium seal with a green Flip-Off sealing.

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:

Global Vet Health S.L.

Marketing authorisation date:

19/12/2012

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 36167/3001

Date of authorisation status change:

28/02/2024

Reference member state:

Ireland

Procedure number:

IE/V/0290/001

Concerned member states:

Bulgaria Croatia Poland Portugal Romania United Kingdom (Northern Ireland)

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