

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

Not
authorised

- Enrofloxacin

Product identification

Medicine name:

Fenoflox 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. 3 day

Subcutaneous use:

-

Cattle

- Meat and offal. 12 day

- Milk. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Expired

Authorised in:

Luxembourg

Package description:

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 250 ml No. of containers in a carton: 20 x 250 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 250 ml No. of containers in a carton: 15 x 250 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 250 ml No. of containers in a carton: 12 x 250 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 250 ml No. of containers in a carton: 10 x 250 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 250 ml No. of containers in a carton: 5 x 250 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 250 ml No. of containers in a carton: 1 x 250 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 100 ml No. of containers in a carton: 20 x 100 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 100 ml No. of containers in a carton: 15 x 100 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 100 ml No. of containers in a carton: 12 x 100 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 100 ml No. of containers in a carton: 10 x 100 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 100 ml No. of containers in a carton: 5 x 100 ml

Container material: Type I Amber Glass Container closure: Grey teflonised chlorobutyl rubber stopper with an aluminium cap. Container colour: Amber Container volume: 100 ml No. of containers in a carton: 1 x 100 ml

Additional information

Entitlement type:

Marketing Authorisation

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

1/03/2010

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Chanelle Pharmaceuticals Manufacturing Ltd

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

V 666/11/01/1048

Date of authorisation status change:

1/03/2010

Reference member state:

Ireland

Procedure number:

IE/V/0223/002

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

Documents

ie-spc-vpa10987-072-002-en.pdf