

Fenoflox 50 mg/ml Solution for Injection for Cattle, Pigs, Dogs and Cats

Authorised

- Enrofloxacin

Product identification

Medicine name:

Fenoflox 50 mg/ml Solution for Injection for Cattle, Pigs, Dogs and Cats

Active substance:

Enrofloxacin

Target species:

Pig
Cattle
Dog
Cat

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Enrofloxacin
50.00 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

Subcutaneous use:

-

Cattle

- 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:

Valid

Authorised in:

Finland

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:

13/01/2011

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Finnish Medicines Agency

Authorisation number:

25273

Date of authorisation status change:

13/01/2011

Reference member state:

Ireland

Procedure number:

IE/V/0223/001

Concerned member states:

Austria Belgium Cyprus Czechia Finland France Germany Hungary Italy
Luxembourg Portugal Slovakia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.