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
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:
Ireland

Package description:

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 250 mlNo. of containers in a carton: 20 x 250 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 250 mlNo. of containers in a carton: 15 x 250 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 250 mlNo. of containers in a carton: 12 x 250 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 250 mlNo. of containers in a carton: 10 x 250 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 250 mlNo. of containers in a carton: 5 x 250 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 250 mlNo. of containers in a carton: 1 x 250 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 100 mlNo. of containers in a carton: 20 x 100 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 100 mlNo. of containers in a carton: 15 x 100 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 100 mlNo. of containers in a carton: 12 x 100 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 100 mlNo. of containers in a carton: 10 x 100 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 100 mlNo. of containers in a carton: 5 x 100 ml

Container material: Type I Amber GlassContainer closure: Grey teflonised chlorobutyl rubber stopper with an aluminium capContainer colour: AmberContainer volume: 100 mlNo. 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:
14/05/2010

Manufacturing sites for batch release:
Labiana Life Sciences S.A.
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/072/001

Date of authorisation status change:
14/05/2010

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