

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

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

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

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**Active substance:**

Enrofloxacin

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**Target species:**

Pig

Cattle

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Pig**

- Meat and offal. 13 day

**Intravenous use:**

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**Cattle**

- Meat and offal. 5 day

- Milk. 3 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 12 day

- Milk. 4 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01MA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Marketing authorisation holder:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Marketing authorisation date:**

17/02/2010

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**Manufacturing sites for batch release:**

Labiana Life Sciences S.A.

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/2817502 5/2010

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**Date of authorisation status change:**

22/12/2021

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0223/002

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**Concerned member states:**

Austria Belgium Finland France Germany Hungary Italy Luxembourg  
Portugal Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 16/01/2026

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