

# Enroxil 100 mg/ml solution for injection for cattle and pigs

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

Enroxil 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:**

Ireland

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**Available in:**

Ireland

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**Package description:**

Cardboard box with 1 amber Type I glass bottle of 100 ml with a grey bromobutyl rubber stopper and aluminium cap.

Cardboard box with 1 amber Type I glass bottle of 50 ml with a grey bromobutyl rubber stopper and aluminium cap.

Cardboard box with 1 amber Type II glass bottle of 50 ml with a grey bromobutyl rubber stopper and aluminium cap.

Cardboard box with 1 amber Type II glass bottle of 100 ml with a grey bromobutyl rubber stopper and aluminium cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

10/07/2009

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10774/002/003

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**Date of authorisation status change:**

10/07/2009

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0422/002

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**Concerned member states:**

Austria Germany Netherlands 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