

# Interflox-100, 100 mg/ml solution for injection for cattle, sheep, goats and pigs

Authorised

- Enrofloxacin

## Product identification

### Medicine name:

Interflox-100, 100 mg/ml solution for injection for cattle, sheep, goats and pigs  
Interflox 100, 100mg/ml, Injekční roztok

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### Active substance:

Enrofloxacin

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### Target species:

Cattle  
Sheep  
Goat  
Pig

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### Route of administration:

Intravenous use  
Subcutaneous use  
Intramuscular 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:**

**Intravenous use:**

• **Cattle**

- Meat and offal. 5 day
- Milk. 3 day

**Subcutaneous use:**

• **Cattle**

- Meat and offal. 12 day
- Milk. 4 day

• **Sheep**

- Meat and offal. 4 day
- Milk. 3 day

• **Goat**

- Meat and offal. 6 day
- Milk. 4 day

**Intramuscular use:**

• **Pig**

- Meat and offal. 13 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:**

Czechia

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**Package description:**

100 ml amber glass bottles (type I) closed with brombutyl rubber stopper and aluminium cap in cardboard box.

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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:**

Interchemie Werken De Adelaar Eesti AS

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**Marketing authorisation date:**

13/05/2019

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**Manufacturing sites for batch release:**

Interchemie Werken De Adelaar Eesti AS

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/030/19-C

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**Date of authorisation status change:**

13/05/2019

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**Reference member state:**

Estonia

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**Procedure number:**

EE/V/0103/001

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**Concerned member states:**

Austria Bulgaria Croatia Cyprus Czechia France Greece Hungary Italy

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

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.

### Labelling

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