

# Quiflor 100 mg/ml solution for injection for cattle and pigs (sows)

Authorised

- Marbofloxacin

## Product identification

**Medicine name:**

Quiflor 100 mg/ml solution for injection for cattle and pigs (sows)

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**Active substance:**

Marbofloxacin

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**Target species:**

Cattle

Pig (sow for reproduction)

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**Route of administration:**

Intravenous use

Subcutaneous use

Intramuscular use

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## Product details

**Active substance and strength:**

Marbofloxacin

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:**

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**Cattle**

- Milk. 36 hour
- Meat and offal. 6 day

**Subcutaneous use:**

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**Cattle**

- Milk. 36 hour
- Meat and offal. 6 day

**Intramuscular use:**

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**Cattle**

- Milk. 36 hour
- Meat and offal. 6 day

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**Pig (sow for reproduction)**

- Meat and offal. 4 day

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

QJ01MA93

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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:**

Slovakia

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**Package description:**

(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)

(ID1) 250 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 250 millilitre(s), closed with Stopfen (bromobutyl rubber)

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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:**

7/07/2011

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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:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/039/DC/11-S

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**Date of authorisation status change:**

7/07/2011

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0302/001

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**Concerned member states:**

Austria Belgium Denmark Greece Italy Lithuania Netherlands Portugal  
Slovakia Spain

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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

Combined File of all Documents

English (PDF)

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