# 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)
Quiflor 100 mg/ml injekčný roztok pre hovädzí dobytok a ošípané (prasnice)

**Active substance:**
Marbofloxacin

**Target species:**
Cattle
Pig (sow for reproduction)

**Route of administration:**
Intravenous use
Subcutaneous use
Intramuscular use

## Product details

**Active substance and strength:**
Marbofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**

**Intravenous use:**

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

**Subcutaneous use:**

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

**Intramuscular use:**

- **Cattle**
  - Milk. 36 hour
  - Meat and offal. 6 day
- **Pig (sow for reproduction)**
  - Meat and offal. 4 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01MA93

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Slovakia

**Package description:**
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (Braunglas) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber`)
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Braunglas) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber`)
(ID1) 250 millilitre(s): unspecified outer container with 1 Vial (Braunglas) with 250 millilitre(s), closed with Stopfen (bromobutyl rubber`)

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
KRKA tovarna zdravil d.d. Novo mesto

**Marketing authorisation date:**
7/07/2011

**Manufacturing sites for batch release:**
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

**Responsible authority:**
Institute For State Control Of Veterinary Biologicals And Medicaments

**Authorisation number:**
96/039/DC/11-S

**Date of authorisation status change:**
7/07/2011

**Reference member state:**
Germany

**Procedure number:**

DE/V/0302/001

**Concerned member states:**
Austria Belgium Denmark Greece Italy Lithuania Netherlands Portugal Slovakia Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents

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