

# MARFLOQUIN 100 MG/ML SOLUTION FOR INJECTION FOR BOVINES AND PIGS (SOWS)

Authorised

- Marbofloxacin

## Product identification

**Medicine name:**

MARFLOQUIN 100 MG/ML SOLUTION FOR INJECTION FOR BOVINES AND PIGS (SOWS)  
Marfloquin 100 mg/ml Injektionslösung für Rinder und Schweine (Sauen)

**Active substance:**

Marbofloxacin

**Target species:**

Cattle

Pig (sow)

**Route of administration:**

Intramuscular use

Subcutaneous use

Intravenous use

## 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:**

**Intramuscular use:**

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**Cattle**

- Milk. 36 hour 2 mg/kg single daily injection, for 3 days
- Meat and offal. 6 day 2 mg/kg single daily injection, for 3 days
- Milk. 72 hour 8 mg/kg single dose
- Meat and offal. 3 day 8 mg/kg single dose

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**Pig (sow)**

- Meat and offal. 4 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 6 day 2 mg/kg single daily injection, for 3 days
- Milk. 36 hour 2 mg/kg single daily injection, for 3 days

**Intravenous use:**

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**Cattle**

- Meat and offal. 6 day

2 mg/kg single daily injection, for 3 days (the first injection may also be given by the intravenous route too)

- Milk. 36 hour

2 mg/kg single daily injection, for 3 days (the first injection may also be given by the intravenous route too)

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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:**

Germany

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**Package description:**

Available only in French

Available only in French

Available only in French

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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:**

7/03/2011

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

Virbac

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

401400.00.00

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**Date of authorisation status change:**

13/06/2016

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**Reference member state:**

France

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**Procedure number:**

FR/V/0223/002

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**Concerned member states:**

Austria Belgium Czechia Germany Greece Hungary Italy Latvia Lithuania  
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

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