

Marfloxin 100 mg/ml solution for injection for cattle and pigs

Authorised

- Marbofloxacin

Product identification

Medicine name:

Marfloxin 100 mg/ml solution for injection for cattle and pigs

Active substance:

Marbofloxacin

Target species:

Cattle

Pig

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous 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:**Intramuscular use:**

-

Cattle

- Meat and offal. 4 day
- Meat and offal. 3 day When administered once at 8 mg/kg
- Milk. 24 hour
- Milk. 72 hour When administered once at 8 mg/kg

-

Pig

- Meat and offal. 2 day

Intravenous use:

-

Cattle

- Meat and offal. 4 day
- Milk. 24 hour

Subcutaneous use:

-

Cattle

- Meat and offal. 4 day
- Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA93

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Available in:

Romania

Package description:

50 ml amber glass vial (Ph. Eur. type II) sealed with a bromobutyl rubber stopper and aluminium closure packaged in an outer carton.

100 ml amber glass vial (Ph. Eur. type II) sealed with a bromobutyl rubber stopper and aluminium closure packaged in an outer carton.

250 ml amber glass vial (Ph. Eur. type II) sealed with a bromobutyl rubber stopper and aluminium closure packaged in an outer carton.

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:

5/02/2012

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

160316

Date of authorisation status change:

3/07/2025

Reference member state:

Ireland

Procedure number:

IE/V/0262/002

Concerned member states:

Bulgaria Poland Romania Slovenia

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

Documents

Combined File of all Documents

English (PDF)

Published on: 6/07/2025

[Download](#)

Summary of Product Characteristics