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

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Authorisation status:
Valid

Authorised in:
Slovenia

Available in:
Slovenia

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:
17/05/2011

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0426/002

Date of authorisation status change:
17/05/2011

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

Summary of Product Characteristics

English (PDF)
Published on: 15/01/2026
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Combined File of all Documents

English (PDF)
Published on: 6/07/2025
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