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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)

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:
Portugal

Package description:
(ID3) 50 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 50 millilitre(s), closed with Stopper (Bromobutyl Rubber)
(ID2) 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 100 millilitre(s), closed with Stopper (Bromobutyl Rubber)
(ID1) 250 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 250 millilitre(s), closed with Stopper (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:
19/05/2011

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
349/01/11DFVPT

Date of authorisation status change:
30/06/2025

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

2401422-paren-20180605.rtf