Quiflor 100 mg/ml solution for injection for cattle and pigs (sows)

Authorised

- Marbofloxacin

Product identification

Medicine name:
QUIFLOR 100MG/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ
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:
Greece

Available in:
Greece

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:
19/01/2012

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

Responsible authority:
National Organization For Medicines

Authorisation number:
86458/10-11-2016/K-0189202

Date of authorisation status change:
30/08/2022

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