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:
Greece

Available in:
Greece

Package description:
(ID1) 250 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 250 millilitre(s), closed with Stopfen (bromobutyl rubber)
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 50 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

Documents

Combined File of all Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

2401422-paren-20180605.rtf