File downloaded on 2026-11-11
Source URL: https://medicines.health.europa.eu/veterinary/en/600000060699

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

Not 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:
Surrendered

Authorised in:
Spain

Package description:
(ID1) 250 millilitre(s): unspecified outer container with 1 Vial (Glass) with 250 millilitre(s), closed with Stopper (bromobutyl rubber`)
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s), closed with Stopper (bromobutyl rubber`)
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 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:
14/12/2011

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2419 ESP

Date of authorisation status change:
20/12/2023

Reference member state:

Germany

Procedure number:
DE/V/0302/001

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

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.

Labelling

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