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# Quiflor 20 mg/ml solution for injection for cattle, pigs and dogs

Not authorised

- Marbofloxacin

## Product identification

**Medicine name:**
Quiflor 20 mg/ml solution for injection for cattle, pigs and dogs

**Active substance:**
Marbofloxacin

**Target species:**
Cattle (pre-ruminant)
Dog
Pig

**Route of administration:**
Intravenous use
Subcutaneous use
Intramuscular use

## Product details

**Active substance and strength:**

Marbofloxacin
20.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**
**Intravenous use:**

- **Cattle (pre-ruminant)**
  - Meat and offal. 6 day

**Subcutaneous use:**

- **Cattle (pre-ruminant)**
  - Meat and offal. 6 day

**Intramuscular use:**

- **Cattle (pre-ruminant)**
  - Meat and offal. 6 day

- **Pig**
  - 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:**
United Kingdom (Northern Ireland)

**Package description:**
(ID4): 1 unspecified outer container with 1 Vial (Glass) with 20 millilitre(s) (20 millilitre(s))
(ID3): 1 unspecified outer container with 1 Vial (Glass) with 250 millilitre(s) (250 millilitre(s))
(ID2): 1 unspecified outer container with 1 Vial (Glass) with 100 millilitre(s) (100 millilitre(s))
(ID1): 1 unspecified outer container with 1 Vial (Glass) with 50 millilitre(s) (50 millilitre(s))

## 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:**
19/12/2011

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

**Responsible authority:**
The Veterinary Medicines Directorate

**Authorisation number:**
Vm01656/4060

**Date of authorisation status change:**
23/05/2022

**Reference member state:**

Germany

**Procedure number:**
DE/V/0301/001

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

## Documents

Summary of Product Characteristics