File downloaded on 2026-11-23
Source URL: https://medicines.health.europa.eu/veterinary/en/600000060783

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:
Netherlands

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:
9/05/2011

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 107423

Date of authorisation status change:
8/02/2022

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

Combined File of all Documents

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

Summary of Product Characteristics