

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