

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

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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:

Netherlands

Package description:

(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 50 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)

(ID1) 250 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 250 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:

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 107424

Date of authorisation status change:

13/09/2022

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

2401422-paren-20180605.rtf