Quiflor S 100 mg/ml solution for injection for cattle

Authorised

Marbofloxacin

Product identification

Medicine name:

Quiflor S 100 mg/ml solution for injection for cattle Quiflor S 100 mg/ml oplossing voor injectie voor runderen

Active substance:

Marbofloxacin

Target species:

Cattle

Route of administration:

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:

Intramuscular use:

- . Cattle
 - Meat and offal. 3 day
 - Milk. 72 hour

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:

(ID2) 250 millilitre(s): unspecified outer container with 1 Vial (Glass) with 250 millilitre(s), closed with Stopper (bromobutyl rubber`) (ID1) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100

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:

9/05/2011

Manufacturing sites for batch release:

Tad Pharma GmbH
Responsible authority: Medicines Evaluation Board
Authorisation number: REG NL 107425
Date of authorisation status change: 3/03/2022
Reference member state: Germany
Procedure number: DE/V/0303/001
Concerned member states: Austria Belgium Denmark Greece Italy Lithuania Netherlands Portugal Slovakia Spain United Kingdom (Northern Ireland)
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Summary of Product Characteristics
Combined File of all Documents

Krka d.d. Novo Mesto

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