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 (Braunglas) with 250 millilitre(s), closed with Stopfen (bromobutyl rubber`)
(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (Braunglas) with 100 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 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

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

Documents

Combined File of all Documents

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