MARFLOQUIN 100 MG/ML SOLUTION FOR INJECTION FOR BOVINES AND PIGS (SOWS)

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARFLOQUIN 100 MG/ML SOLUTION FOR INJECTION FOR BOVINES AND PIGS (SOWS)
Quiflox, 100mg/ml, Injekční roztok

Active substance:
Marbofloxacin

Target species:
Cattle
Pig (sow)

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous 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**
  - Milk. 36 hour — 2 mg/kg single daily injection, for 3 days
  - Meat and offal. 6 day — 2 mg/kg single daily injection, for 3 days
  - Milk. 72 hour — 8 mg/kg single dose
  - Meat and offal. 3 day — 8 mg/kg single dose
- **Pig (sow)**
  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 6 day — 2 mg/kg single daily injection, for 3 days
  - Milk. 36 hour — 2 mg/kg single daily injection, for 3 days

Intravenous use:

- **Cattle**
  - Meat and offal. 6 day

2 mg/kg single daily injection, for 3 days (the first injection may also be given by the intravenous route too)

- Milk. 36 hour

2 mg/kg single daily injection, for 3 days (the first injection may also be given by the intravenous route too)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
Box of one 50 ml solution bottle for injection
Box of one 250 ml solution bottle for injection
Box of one 100 ml solution bottle for injection

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:
6/09/2011

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
Virbac

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/077/11-C

Date of authorisation status change:
7/06/2016

Reference member state:
France

Procedure number:
FR/V/0223/002

Concerned member states:
Austria Belgium Czechia Germany Greece Hungary Italy Latvia Lithuania Portugal Slovakia Spain

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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