

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)
Marfloquin 100 mg/ml Injektionslösung für Rinder und Schweine (Sauen)

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:

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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

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Pig (sow)

- Meat and offal. 4 day

Subcutaneous use:

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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:

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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:

Germany

Package description:

Available only in French

Available only in French

Available only in French

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:

7/03/2011

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Virbac

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401400.00.00

Date of authorisation status change:

13/06/2016

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

Combined File of all Documents

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