MARBOSYVA 100 mg/ml solution for injection for cattle and pigs.

Authorised

- Marbofloxacin

Product identification

Medicine name:
MARBOSYVA 100 mg/ml solution for injection for cattle and pigs.
Marbosyva 100 mg/ml Injektionslösung für Rinder und Schweine

Active substance:
Marbofloxacin

Target species:
Cattle
Pig (female)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous 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. no withdrawal period

Meat and offal: Intramuscular use (8 mg/kg single dose): 3 days / Intramuscular or subcutaneous use (2 mg/kg single daily injection, for 3-5 days): 6 days

- **Pig (female)**
  - Meat and offal. 4 day

- **Cattle**
  - Milk. no withdrawal period

Milk: IM (single dose) 72 hours / IM or SC 36 Hours

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period

Meat and offal: Intramuscular use (8 mg/kg single dose): 3 days / Intramuscular or subcutaneous use (2 mg/kg single daily injection, for 3-5 days): 6 days

- **Cattle**
  - Milk. no withdrawal period

Milk: IM (single dose) 72 hours / IM or SC 36 Hours

Subcutaneous use:

- **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Intramuscular use (8 mg/kg single dose): 3 days / Intramuscular or subcutaneous use (2 mg/kg single daily injection, for 3-5 days): 6 days

- **Cattle**
  - Milk. no withdrawal period

Milk: IM (single dose) 72 hours / IM or SC 36 Hours

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German
Available only in German
Available only in German

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:
Laboratorios Syva S.A.

Marketing authorisation date:
13/11/2012

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-01130

Date of authorisation status change:
13/11/2012

Reference member state:
Spain

Procedure number:
ES/V/0188/001

Concerned member states:
Austria Bulgaria Germany Hungary Ireland Italy Poland Portugal Romania United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

English (PDF)
Published on: 22/12/2023

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Summary of Product Characteristics

English (PDF)
Published on: 22/12/2023
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Labelling

English (PDF)
Published on: 22/12/2023
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Combined File of all Documents

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