Marbosol 100 mg/ml solution for injection for cattle and pigs

Authorised

- Marbofloxacin

Product identification

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

Active substance:
Marbofloxacin

Target species:
Cattle
Pig

Route of administration:
Intravenous use
Subcutaneous use
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:

Intravenous use:

- **Cattle**

- Meat and offal. 6 day

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days) (IV/SC/IM)

- Milk. 36 hour

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days) (IV/SC/IM)

Subcutaneous use:

- **Cattle**

- Meat and offal. 6 day

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days) (IV/SC/IM)

- Milk. 36 hour

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days) (IV/SC/IM)

Intramuscular use:

- **Cattle**

- Meat and offal. 6 day

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days) (IV/SC/IM)

- Milk. 72 hour Single dose (8 mg/kg) (IM)

- Milk. 36 hour

Multiple dose (2 mg/kg single daily injection, for 3 to 5 days) (IV/SC/IM)

- Meat and offal. 3 day Single dose (8 mg/kg) (IM)

- **Pig**

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s)
(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)

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:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
19/01/2013

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401633.00.00

Date of authorisation status change:
10/08/2018

Reference member state:
Germany

Procedure number:
DE/V/0175/002

Concerned member states:
Hungary Spain

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

Documents

Summary of Product Characteristics

English (RTF)
Published on: 18/12/2024
Download

2401633-paren-20170418.pdf

Source URL: *https://medicines.health.europa.eu/veterinary/600000060415*