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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

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**
  - Milk. 36 hour Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)
  - Meat and offal. 6 day Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

Subcutaneous use:

- **Cattle**
  - Meat and offal. 6 day Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)
  - Milk. 36 hour Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

Intramuscular use:

- **Pig**
  - Meat and offal. 4 day
- **Cattle**
  - Meat and offal. 3 day Single dose (8 mg/kg)
  - Meat and offal. 6 day Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)
  - Milk. 72 hour Single dose (8 mg/kg)
  - Milk. 36 hour Multiple dose (2 mg/kg single daily injection, for 3 to 5 days)

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:
(ID1) 50 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 50 millilitre(s)
(ID2) 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 100 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

Combined File of all Documents

English (PDF)
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