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Marbodug 100 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Marbofloxacin

Product identification

Medicine name:
Marbodug 100 mg/ml Solution for Injection for Cattle and Pigs

Active substance:
Marbofloxacin

Target species:
Cattle
Pig

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. 3 day — 8mg/kg on a single occasion (IM)
  - Meat and offal. 6 day — 2 mg/kg for 3 to 5 days
  - Milk. 36 hour — 2 mg/kg for 3 to 5 days
  - Milk. 72 hour — 8mg/kg on a single occasion (IM)
- **Pig**
  - Meat and offal. 4 day

Intravenous use:

- **Cattle**
  - Meat and offal. 6 day
  - Milk. 36 hour

Subcutaneous use:

- **Cattle**
  - Meat and offal. 6 day — 2mg/kg for 3 to 5 days
  - Milk. 36 hour — 2mg/kg 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:
Packaged in Amber type II glass vial of 250ml.Vial is closed with a fluorinated bromobutyl rubber stopper oversealed with an aluminium cap. Each vial is packaged in a cardboard box.
Packaged in Amber type II glass vial of 100ml.Vial is closed with a fluorinated bromobutyl rubber stopper oversealed with an aluminium cap. Each vial is packaged in a cardboard box.
Packaged in Amber type II glass vial of 50ml.Vial is closed with a fluorinated bromobutyl rubber stopper oversealed with an aluminium cap. Each vial is packaged in a cardboard box.
Packaged in Amber type II glass vial of 20ml.Vial is closed with a fluorinated bromobutyl rubber stopper oversealed with an aluminium cap. Each vial is packaged in a cardboard box.
Packaged in Amber type II glass vial of 10ml.Vial is closed with a fluorinated bromobutyl rubber stopper oversealed with an aluminium cap. Each vial is packaged in a cardboard box.

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:
Emdoka

Marketing authorisation date:
4/11/2012

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401637.00.00

Date of authorisation status change:
22/02/2018

Reference member state:
Ireland

Procedure number:
IE/V/0457/002

Concerned member states:
Austria Belgium Germany Luxembourg Netherlands Portugal Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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