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
Odimar 100 mg/ml Injektionslösung für Rinder und Schweine

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:
Austria

Available in:
Austria

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

Marketing authorisation date:
28/11/2012

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-01134

Date of authorisation status change:
28/11/2012

Reference member state:
Ireland

Procedure number:
IE/V/0457/002

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

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 15/12/2024
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Package Leaflet

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Labelling

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