

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:

Portugal

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:

23/10/2012

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

605/02/12DFVPT

Date of authorisation status change:

20/12/2024

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

Combined File of all Documents

English (PDF)

Published on: 15/12/2024

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