

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

Authorised

This information is not available for this product.

Product identification

Medicine name:

Marbodug 100 mg/ml Solution for Injection for Cattle and Pigs
Odimar 100 mg/ml Oplossing voor injectie
Odimar 100 mg/ml Solution injectable
Odimar 100 mg/ml Injektionslösung

Active substance:

This information is not available for this product.

Target species:

Cattle
Pig

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

This information is not available for this product.

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
- Milk. 36 hour
- 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:

Belgium

Available in:

Belgium

Package description:

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.

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 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 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 250ml. 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:

1/03/2013

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V434752

Date of authorisation status change:

1/03/2013

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: 11/02/2022

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Package Leaflet

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