Marbonor 100 mg/ml Solution for Injection for cattle and pig

Authorised

- Marbofloxacin

Product identification

Medicine name:
Marbonor 100 mg/ml Solution for Injection for cattle and pig
Marbonor 100mg/ml ενέσιμο διάλυμα για Βοοειδή και Χοίρους

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. 6 day
  - Milk. 36 hour
- **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
  - Milk. 36 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
The product is packaged in 60 ml amber co-ex plastic (polypropylene) vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 500 ml amber type II glass vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 250 ml amber type II glass vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 100 ml amber type II glass vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 50 ml amber type II glass vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 20 ml amber type II glass vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 100 ml amber co-ex plastic (polypropylene) vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 250 ml amber co-ex plastic (polypropylene) vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.
The product is packaged in 500 ml amber co-ex plastic (polypropylene) vial.Vial is closed with chlorobutyl rubber stopper sealed with aluminium cap.

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
11/06/2013

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
National Organization For Medicines

Authorisation number:
21012/02-03-2018/K-0196302

Date of authorisation status change:
10/02/2020

Reference member state:
Ireland

Procedure number:
IE/V/0296/001

Concerned member states:
Cyprus Czechia Estonia Finland Greece Hungary Latvia Romania Slovakia Slovenia 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: 28/01/2022
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Package Leaflet

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Combined File of all Documents