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

Not authorised

- Marbofloxacin

Product identification

Medicine name:
MARBONOR 100 mg/ml, soluție injectabilă pentru bovine și porcine
Marbonor 100 mg/ml Solution for Injection for cattle and pig

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:
Surrendered

Authorised in:
Romania

Package description:
The product is packaged in 60 ml amber co-ex plastic (polypropylene) vials. The vials are closed with chlorobutyl rubber stoppers sealed with aluminium caps.
The product is packaged in 500 ml amber type II glass vials and amber co-ex plastic (polypropylene) vials. The vials are closed with chlorobutyl rubber stoppers sealed with aluminium caps.
The product is packaged in 250 ml amber type II glass vials and amber co-ex plastic (polypropylene) vials. The vials are closed with chlorobutyl rubber stoppers sealed with aluminium caps.
The product is packaged in 100 ml amber type II glass vials and amber co-ex plastic (polypropylene) vials. The vials are closed with chlorobutyl rubber stoppers sealed with aluminium caps.
The product is packaged in 50 ml amber type II glass vials. The vials are closed with chlorobutyl rubber stoppers sealed with aluminium caps.
The product is packaged in 20 ml amber type II glass vials. The vials are closed with chlorobutyl rubber stoppers sealed with aluminium caps.

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:
16/07/2013

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
190097

Date of authorisation status change:
27/03/2022

Reference member state:
Ireland

Procedure number:
IE/V/0296/001

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

Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics