Marbocyl 20 mg/ml Solution for Injection

Authorised

- Marbofloxacin

Product identification

Medicine name:
Marbocyl 20 mg/ml Solution for Injection

Active substance:
Marbofloxacin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Marbofloxacin
20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 4 day

- **Pig**
  - Meat and offal. 2 day

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA93

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
Marbocyl 2% injection is packaged in amber type II glass vial of 10 ml.The vial is closed with a chlorobutyl rubber stopper and oversealed with aluminium cap.Each vial is packaged in a cardboard box.
Marbocyl 2% injection is packaged in amber type II glass vial of 20 ml.The vial is closed with a chlorobutyl rubber stopper and oversealed with aluminium cap.Each vial is packaged in a cardboard box.
Marbocyl 2% injection is packaged in amber type II glass vial of 50 ml.The vial is closed with a chlorobutyl rubber stopper and oversealed with aluminium cap.Each vial is packaged in a cardboard box.
Marbocyl 2% injection is packaged in amber type II glass vial of 100 ml.The vial is closed with a chlorobutyl rubber stopper and oversealed with aluminium cap.Each vial is packaged in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Vetoquinol Ireland Limited

Marketing authorisation date:
18/09/1998

Manufacturing sites for batch release:
Vetoquinol S.A.
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10983/032/001

Date of authorisation status change:
18/09/1998

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

Documents

Summary of Product Characteristics