

Marbocyl 100 mg/ml Solution for Injection

Authorised

- Marbofloxacin

Product identification

Medicine name:

Marbocyl 100 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

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. 4 day
- Milk. 24 hour

-

Pig

- Meat and offal. 2 day

Intravenous use:

-

Cattle

- Meat and offal. 4 day
- Milk. 24 hour

Subcutaneous use:

-

Cattle

- Meat and offal. 4 day
- Milk. 24 hour

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 10% is packaged in amber type II glass vial of 10 ml. The vial is closed with a chlorobutyl rubber stopper oversealed with aluminium cap. Each vial is packaged in a cardboard box.

Marbocyl 10% is packaged in amber type II glass vial of 20 ml. The vial is closed with a chlorobutyl rubber stopper oversealed with aluminium cap. Each vial is packaged in a cardboard box.

Marbocyl 10% is packaged in amber type II glass vial of 50 ml. The vial is closed with a chlorobutyl rubber stopper oversealed with aluminium cap. Each vial is packaged in a cardboard box.

Marbocyl 10% is packaged in amber type II glass vial of 100 ml. The vial is closed with a chlorobutyl rubber stopper oversealed with aluminium cap. Each vial is packaged in a cardboard box.

Marbocyl 10% is packaged in amber type II glass vial of 250 ml. The vial is closed with a chlorobutyl rubber stopper 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:

31/10/1997

Manufacturing sites for batch release:

Vetoquinol S.A.

Vetoquinol Biowet Sp. z o.o.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10983/034/001

Date of authorisation status change:

31/10/1997

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

Documents

Summary of Product Characteristics