

# MARBOCYL 100 MG/ML, SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Marbofloxacin

## Product identification

**Medicine name:**

MARBOCYL 100 MG/ML, SOLUTION FOR INJECTION FOR CATTLE AND PIGS

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**Active substance:**

Marbofloxacin

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Marbofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

- Meat and offal. 6 day

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

- Milk. 36 hour

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Meat and offal. 6 day

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Milk. 72 hour

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Meat and offal. 3 day

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Milk. 36 hour

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

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**Pig**

- Meat and offal. 4 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 6 day

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

- Milk. 36 hour

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Meat and offal. 6 day

Indication : Mastitis Dosage : 2mg/kg for 3 days (IV/IM/SC)

- Milk. 72 hour

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Meat and offal. 3 day

Indication : Respiratory Dosage : 8 mg/kg on a single occasion (IM)

- Milk. 36 hour

Indication : Respiratory Dosage : 2 m/kg for 3 to 5 days (IV/IM/SC)

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01MA93

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Available in:**

Italy

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetoquinol S.A.

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**Marketing authorisation date:**

17/10/1998

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**Manufacturing sites for batch release:**

Vetoquinol Biowet Sp. z o.o.

Vetoquinol S.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

102602

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**Date of authorisation status change:**

17/10/1998

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**Reference member state:**

France

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**Procedure number:**

FR/V/0107/001

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**Concerned member states:**

Austria Belgium Germany Greece Italy Luxembourg Portugal Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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