

# BOFLOX 100 mg/ml

Authorised

- Marbofloxacin

## Product identification

**Medicine name:**

BOFLOX 100 mg/ml

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**Active substance:**

Marbofloxacin

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**Target species:**

Cattle

Pig (sow)

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**Route of administration:**

Intramuscular use

Intravenous 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. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

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#### **Pig (sow)**

- Meat and offal. 4 day

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#### **Cattle**

- Milk. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

### **Intravenous use:**

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#### **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

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#### **Cattle**

- Milk. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

### **Subcutaneous use:**

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#### **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

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### **Cattle**

- Milk. no withdrawal period

Meat and offal: 6 days 2 mg (IV/SC/IM) /3 days 8 mg (IM); Milk:36 hours 2 mg (IV/SC/IM) / 72 hours 8 mg (IM)

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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:**

Germany

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### **Available in:**

Germany

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### **Package description:**

cardboard box containing 12 vials of 250 ml

cardboard box containing 12 vials of 100 ml

cardboard box containing 10 vials of 250 ml

cardboard box containing 10 vials of 100 ml

cardboard box containing 6 vials of 250 ml

cardboard box containing 6 vials of 100 ml

cardboard box containing 1 vial of 250 ml

cardboard box containing 1 vial of 100 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Industrial Veterinaria S.A.

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**Marketing authorisation date:**

6/03/2013

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**Manufacturing sites for batch release:**

Industrial Veterinaria S.A.

KELA Kempisch Laboratorium Kela Laboratoria

aniMedica GmbH

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

401769.00.00

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**Date of authorisation status change:**

10/08/2018

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0190/001

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**Concerned member states:**

Belgium Bulgaria Cyprus Czechia France Germany Greece Hungary Ireland

Italy Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia

United Kingdom (Northern Ireland)

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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

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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