

# Paramectin 10 mg/ml Solution for Injection

Authorised

- Ivermectin

## Product identification

**Medicine name:**

Paramectin 10 mg/ml Solution for Injection

Paramectin 10 mg/ml Solução Injetável para bovinos e suínos

**Active substance:**

Ivermectin

**Target species:**

Cattle

Pig

**Route of administration:**

Subcutaneous use

## Product details

**Active substance and strength:**

Ivermectin

10.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 49 day

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**Pig**

- Meat and offal. 18 day

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

QP54AA01

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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:**

Portugal

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**Available in:**

Portugal

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**Package description:**

The product will be supplied in 50 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 100 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 250 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 500 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 1 litre volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

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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:**

Norbrook Laboratories Limited

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**Marketing authorisation date:**

12/09/2006

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**Manufacturing sites for batch release:**

Norbrook Manufacturing Ltd

Norbrook Laboratories Limited

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

51396

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**Date of authorisation status change:**

27/01/2025

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0119/001

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**Concerned member states:**

Greece Italy 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

Summary of Product Characteristics

Combined File of all Documents

English (PDF)

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