

# Qualimec 10 mg/ml Solution for Injection.

Authorised

- Ivermectin

## Product identification

**Medicine name:**

Qualimec 10 mg/ml Solution for Injection.  
Ivermax 10 mg/ml oplossing voor injectie

**Active substance:**

Ivermectin

**Target species:**

Cattle  
Sheep  
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

**Withdrawal period by route of administration:****Subcutaneous use:****• Cattle**

- Meat and offal. 49 day

**• Sheep**

- Meat and offal. 42 day

**• Pig**

- Meat and offal. 28 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:**

Netherlands

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**Package description:**

Clear PET multidose container with bromobutyl rubber stopper and aluminium cap.

Pack size 50 ml

HDPE multidose container with bromobutyl rubber stopper and aluminium cap. Pack size: 200 ml

Clear PET multidose container with bromobutyl rubber stopper and aluminium cap.

Pack size 500 ml

HDPE multidose container with bromobutyl rubber stopper and aluminium cap. Pack size: 500 ml.

HDPE multidose container with bromobutyl rubber stopper and aluminium cap. Pack size: 50 ml

Clear PET multidose container with bromobutyl rubber stopper and aluminium cap.

Pack size 250 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Eco Animal Health Europe Limited

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**Marketing authorisation date:**

12/01/2007

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

Divasa Farmavic S.A.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 10446

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**Date of authorisation status change:**

26/01/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0145/001

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**Concerned member states:**

France Germany Greece Hungary Italy Luxembourg Netherlands Portugal  
Spain 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

Combined File of all Documents

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Summary of Product Characteristics

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