

# Ecomectin 10 mg/ml Solution for Injection

Authorised

- Ivermectin

## Product identification

**Medicine name:**

Ecomectin 10 mg/ml Solution for Injection

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**Active substance:**

Ivermectin

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**Target species:**

Cattle

Sheep

Pig

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Ivermectin

10.00 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:**

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 42 day

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**Sheep**

- Meat and offal. 42 day

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**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:**

Medicinal product subject to medical prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Package description:**

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

Pack size 250 ml

HDPE 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: 500 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: 200 ml.

Clear PET multidose container with bromobutyl rubber stopper and aluminium cap.  
Pack size 50 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:**

2/03/2010

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

Divasa Farmavic S.A.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

14796/03-03-2010/K-0150301

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**Date of authorisation status change:**

14/07/2021

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0144/001

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**Concerned member states:**

Austria Belgium Estonia France Germany Greece Italy Latvia Lithuania  
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

Summary of Product Characteristics

Combined File of all Documents