

# KEYTIL 300 mg/ml + 90 mg/ml solution for injection

Not  
authorised

- Tilmicosin
- Ketoprofen

## Product identification

**Medicine name:**

KEYTIL 300 mg/ml + 90 mg/ml solution for injection

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**Active substance:**

Tilmicosin

Ketoprofen

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Tilmicosin

300.00 milligram(s) / 1.00 millilitre(s)

Ketoprofen

90.00 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. 93 day
- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

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

QJ01FA99

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Germany

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**Package description:**

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

box containing 1 vial of 50 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Fixed combination application (Article 13b of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetpharma Animal Health S.L.

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**Marketing authorisation date:**

12/12/2018

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**Manufacturing sites for batch release:**

Laboratorios Maymo S.A.U.

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

402457.00.00

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**Date of authorisation status change:**

17/12/2025

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0280/001

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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

English (PDF)

Published on: 1/02/2024

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## Package Leaflet

English (PDF)

Published on: 1/02/2024

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## Labelling

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