

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

Active substance:

Tilmicosin

Ketoprofen

Target species:

Cattle

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Tilmicosin

300.00 milligram(s) / 1.00 millilitre(s)

Ketoprofen

90.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Belgium

Package description:

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

box containing 1 vial of 50 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Vetpharma Animal Health S.L.

Marketing authorisation date:

5/02/2019

Manufacturing sites for batch release:

Laboratorios Maymo S.A.U.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V538862

Date of authorisation status change:

18/12/2025

Reference member state:

Spain

Procedure number:

ES/V/0280/001

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 1/02/2024

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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