

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

Authorised

- Tilmicosin
- Ketoprofen

Product identification

Medicine name:

Tilkomay 300 mg/ml + 90 mg/ml solution for injection
TILKOMAY 300 mg/ml + 90 mg/ml SOLUCION INYECTABLE

Active substance:

Tilmicosin
Ketoprofen

Target species:

Cattle (calf)

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:

-

Cattle (calf)

- 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:

Valid

Authorised in:

Spain

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:

Laboratorios Maymo S.A.U.

Marketing authorisation date:

22/01/2019

Manufacturing sites for batch release:

Laboratorios Maymo S.A.U.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3734 ESP

Date of authorisation status change:

22/01/2019

Reference member state:

Spain

Procedure number:

ES/V/0279/001

Concerned member states:

Hungary Italy Poland Portugal Romania

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 12/04/2023

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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