

TILMINJECT 300 mg/ml solution for injection for cattle and ovine

Authorised

- Tilmicosin

Product identification

Medicine name:

TILMINJECT 300 mg/ml solution for injection for cattle and ovine

Active substance:

Tilmicosin

Target species:

Cattle

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Tilmicosin

300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

-

Cattle

- Meat and offal. 70 day

- Milk. 36 day

-

Sheep

- Meat and offal. 42 day

- Milk. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Cyprus

Package description:

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratorios Calier S.A.

Marketing authorisation date:

11/02/2021

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00813V

Date of authorisation status change:

11/02/2021

Reference member state:

Spain

Procedure number:

ES/V/0333/001

Concerned member states:

Croatia Cyprus Czechia Germany Greece 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: 22/12/2023

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Package Leaflet

Labelling

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