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TILMI-kel 300 mg/ml solution for injection for cattle

Not authorised

- Tilmicosin

Product identification

Medicine name:
TILMI-KEL 300MG/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ
TILMI-kel 300 mg/ml solution for injection for cattle

Active substance:
Tilmicosin

Target species:
Sheep
Cattle

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:

- **Sheep**
  - Meat and offal. 42 day
  - Milk. 18 day
- **Cattle**
  - Meat and offal. 70 day
  - Milk. 36 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Greece

Package description:

box containing 12 vials of 250 ml
box containing 12 vials of 100 ml
box containing 12 vials of 50 ml
box containing 10 vials of 250 ml
box containing 10 vials of 100 ml
box containing 10 vials of 50 ml
box containing 6 vials of 250 ml
box containing 6 vials of 100 ml
box containing 6 vials of 50 ml
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:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Kela Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
24/02/2010

Manufacturing sites for batch release:
Industrial Veterinaria S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
58055/08-09-2015/K-0181001

Date of authorisation status change:
28/12/2022

Reference member state:
Spain

Procedure number:
ES/V/0144/001

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