

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

Not
authorised

- Tilmicosin

Product identification

Medicine name:

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:**

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Sheep

- Meat and offal. 42 day

- Milk. 18 day

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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:

Hungary

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:

14/10/2009

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

2626/X/09 MgSzH ÁTI

Date of authorisation status change:

6/12/2022

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