Tilmovet 300 mg/ml Solution for Injection for cattle and sheep

Authorised

- Tilmicosin

Product identification

Medicine name:
Tilmovet 300 mg/ml Solution for Injection for cattle and sheep

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

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Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

100 ml Type II amber glass vial sealed with Type I bromobutyl stopper and aluminium cap, supplied in cardboard box. One vial per box.
50 ml Type II amber glass vial sealed with Type I bromobutyl stopper and aluminium cap, supplied in cardboard box. One vial per box.
25 ml Type I amber glass vial sealed with Type I bromobutyl stopper and aluminium cap, supplied in cardboard box. One vial per box.
250 ml Type II amber glass vial sealed with Type I bromobutyl stopper and aluminium cap, supplied in cardboard box. One vial per box.

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:
HuVepharma

Marketing authorisation date:
3/05/2019

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402525.00.00

Date of authorisation status change:
3/05/2019

Reference member state:
Ireland

Procedure number:
IE/V/0405/001

Concerned member states:
Belgium Bulgaria France Germany Italy Netherlands Poland Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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