

# MICOTIL 300 MG/ML SOLUTION INJECTABLE

Authorised

- Tilmicosin

## Product identification

**Medicine name:**

MICOTIL 300 MG/ML SOLUTION INJECTABLE

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**Active substance:**

Tilmicosin

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**Target species:**

Cattle

Sheep

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Tilmicosin

300.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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

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**Cattle**

- Meat and offal. 70 day
- Milk. 36 day

Si le produit est administré aux vaches pendant la période de tarissement ou aux génisses laitières gestantes (conformément à la rubrique « Utilisation en cas de gestation, de lactation ou de ponte » ci-dessus), le lait ne doit pas être utilisé pour la consommation humaine jusqu'à 36 jours après le vêlage.

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**Sheep**

- Meat and offal. 42 day
- Milk. 18 day

Si le produit est administré aux brebis pendant la période de tarissement ou aux brebis gestantes (conformément à la rubrique « Utilisation en cas de gestation, de lactation ou de ponte » ci-dessus), le lait ne doit pas être utilisé pour la consommation humaine jusqu'à 18 jours après l'agnelage.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01FA91

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Available in:**

France

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Elanco GmbH

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**Marketing authorisation date:**

4/08/1994

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**Manufacturing sites for batch release:**

Elanco France S.A.S.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/6240822 8/1994

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**Date of authorisation status change:**

4/08/2009

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.