

# Tilmovet 250 mg/ml Concentrate for oral solution

Authorised

- Tilmicosin

## Product identification

**Medicine name:**

Tilmovet 250 mg/ml Concentrate for oral solution

---

**Active substance:**

Tilmicosin

---

**Target species:**

Pig  
Turkey  
Chicken  
Cattle (calf)

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Tilmicosin  
250.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Concentrate for oral solution

---

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

- 

**Pig**

- Meat and offal. 14 day      Chickens: 12 days Turkeys: 19 days

- 

**Turkey**

- Meat and offal. 19 day

Eggs: Not authorised for use in birds producing eggs for human consumption.

- 

**Chicken**

- Meat and offal. 12 day

Eggs: Not authorised for use in birds producing eggs for human consumption.

- 

**Cattle (calf)**

- Meat and offal. 42 day

---

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

QJ01FA91

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Bulgaria

---

**Package description:**

High density polyethylene (HDPE) bottle with a tamper-evident screw closure made of polypropylene (PP)

White high density polyethylene bottle with white polypropylene or high density polyethylene, tamper-evident cap

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

---

**Marketing authorisation holder:**

HuVepharma

---

**Marketing authorisation date:**

19/11/2008

---

**Manufacturing sites for batch release:**

Biovet AD

---

**Responsible authority:**

Bulgarian Food Safety Authority

---

**Authorisation number:**

0022-2079

---

**Date of authorisation status change:**

19/11/2008

---

**Reference member state:**

Belgium

---

**Procedure number:**

BE/V/0016/001

---

**Concerned member states:**

Austria Bulgaria Czechia Denmark France Greece Hungary Ireland Italy

Netherlands Poland Portugal Romania Spain  
United Kingdom (Northern Ireland)

---

**Generic of:**  
600000080058

---

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

English (PDF)  
Published on: 13/03/2026  
[Download](#)

### Package Leaflet

English (PDF)  
Published on: 13/03/2026  
[Download](#)

### Labelling

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

Combined File of all Documents