

# TYLMASIN 200 MG/ML SOLUTION FOR INJECTION FOR CATTLE, SHEEP, GOATS AND PIGS

Authorised

- Tylosin

## Product identification

**Medicine name:**

TYLMASIN 200 MG/ML SOLUTION FOR INJECTION FOR CATTLE, SHEEP, GOATS AND PIGS

Tylmasin 200 mg/ml solutie injectabila pentru bovine, ovine, caprine si porcine

**Active substance:**

Tylosin

**Target species:**

Cattle

Pig

Sheep

Goat

**Route of administration:**

Intramuscular use

Intravenous use

## Product details

**Active substance and strength:**

Tylosin

200000.00 international unit(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for injection

---

**Withdrawal period by route of administration:**

**Intramuscular use:**

- 

**Cattle**

- Meat and offal. 28 day
- Milk. 108 hour

- 

**Pig**

- Meat and offal. 16 day

- 

**Sheep**

- Meat and offal. 42 day
- Milk. 108 hour

- 

**Goat**

- Meat and offal. 42 day
- Milk. 108 hour

**Intravenous use:**

- 

**Cattle**

- Meat and offal. 28 day
- Milk. 108 hour

---

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

QJ01FA90

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Romania

---

**Package description:**

Available only in French

Available only in French

Available only in French

---

## 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:**

Biovet AD

---

**Marketing authorisation date:**

25/02/2013

---

**Manufacturing sites for batch release:**

Biovet AD

---

**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

---

**Authorisation number:**

180087

---

**Date of authorisation status change:**

21/12/2020

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0240/001

---

**Concerned member states:**

Austria Bulgaria Denmark Germany Greece Hungary Ireland Italy Poland  
Portugal Romania Spain

---

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

## Documents

Combined File of all Documents

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

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000040439>