

# Baycox Multi 50 mg/ml oral suspension for Cattle, Pigs and Sheep

Authorised

- Toltrazuril

## Product identification

**Medicine name:**

Baycox Multi 50 mg/ml oral suspension for Cattle, Pigs and Sheep

Baycox Multi 50 mg/ml peroralna suspenzija za govedo, prašiče in ovce

**Active substance:**

Toltrazuril

**Target species:**

Cattle

Sheep

Pig

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Toltrazuril

50.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Oral suspension

---

**Withdrawal period by route of administration:**

**Oral use:**

- 

**Cattle**

- Meat and offal. 63 day

- 

**Sheep**

- Meat and offal. 42 day

- 

**Pig**

- Meat and offal. 77 day

---

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

QP51BC01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Slovenia

---

**Available in:**

Slovenia

---

**Package description:**

Available only in Slovenian

Available only in Slovenian

Available only in Slovenian

---

## 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:**

Elanco Animal Health GmbH

---

**Marketing authorisation date:**

25/01/2019

---

**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

---

**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

---

**Authorisation number:**

DC/V/0029/004

---

**Date of authorisation status change:**

25/01/2019

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0360/001

---

**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Italy Latvia Lithuania  
Luxembourg Netherlands Norway Poland Portugal Romania Slovakia  
Slovenia Spain Sweden United Kingdom (Northern Ireland)

---

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: 6/07/2025

[Download](#)

### Package Leaflet

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

### Labelling

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

### Combined File of all Documents