

# Tullavis 100 mg/ml solution for injection for cattle, pigs and sheep

Authorised

- Tulathromycin

## Product identification

**Medicine name:**

Tullavis 100 mg/ml solution for injection for cattle, pigs and sheep

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**Active substance:**

Tulathromycin

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**Target species:**

Sheep

Pig

Cattle

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Tulathromycin

100.00 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:**

**Intramuscular use:**

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**Sheep**

- Meat and offal. 16 day

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**Pig**

- Meat and offal. 13 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 22 day

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

QJ01FA94

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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:**

Denmark

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**Available in:**

Denmark

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**Package description:**

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 20 ml.

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 50 ml.

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 100 ml.

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 250 ml.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Industrial Veterinaria S.A.

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**Marketing authorisation date:**

13/10/2020

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**Manufacturing sites for batch release:**

aniMedica Herstellungs GmbH

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

63203

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**Date of authorisation status change:**

13/10/2020

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0893/001

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**Concerned member states:**

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

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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

English (PDF)

Published on: 21/12/2023

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Combined File of all Documents

eu-PUAR-esv0358001-dcp-tulavis-100-mg-mlsolution-for-injection-for-cattle--pigs-  
and-sheep-en.pdf