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

Authorised

- Tulathromycin

## Product identification

**Medicine name:**
DAXTON 100 mg/ml solution for injection for cattle, pigs and sheep
Daxton 100 mg/ml solution injectable pour bovins, porcins et ovins
Daxton 100 mg/ml oplossing voor injectie voor rund, varken en schaap
Daxton 100 mg/ml Injektionslösung für Rinder, Schweine und Schafe

**Active substance:**
Tulathromycin

**Target species:**
Cattle
Pig
Sheep

**Route of administration:**
Subcutaneous use
Intramuscular use

## 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:**
**Subcutaneous use:**

- **Cattle**
  - Meat and offal. 22 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption

**Intramuscular use:**

- **Pig**
  - Meat and offal. 13 day
- **Sheep**
  - Meat and offal. 16 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption

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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

**Authorised in:**
Belgium

**Package description:**
box with 1 vial 100 ml
box with 1 vial 250 ml

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 18 of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Cenavisa S.L.

**Marketing authorisation date:**
7/11/2024

**Manufacturing sites for batch release:**
Cenavisa S.L.

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
BE-V663457

**Date of authorisation status change:**
7/11/2024

**Reference member state:**
Spain

**Procedure number:**
ES/V/0428/001

**Concerned member states:**
Belgium Bulgaria Croatia Czechia Denmark Estonia Greece Hungary Italy Latvia Lithuania Poland Romania

**Generic of:**
600000003851
600000004232

To consult adverse reactions on veterinary medicinal products please go to 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.

Labelling

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

Package Leaflet

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

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