

# Draxxin 100 mg/ml - Solution for injection (cattle, pig, sheep)

Authorised

- Tulathromycin

## Product identification

**Medicine name:**

Draxxin 100 mg/ml - Solution for injection (cattle, pig, sheep)

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**Active substance:**

Tulathromycin

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**Target species:**

Cattle

Sheep

Pig

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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 Vial

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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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**Cattle**

- Meat and offal. 22 day 22 days

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**Sheep**

- Meat and offal. 16 day 16 days

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**Pig**

- Meat and offal. 13 day 13 days

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 22 day 22 days

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**Sheep**

- Meat and offal. 16 day 16 days

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**Pig**

- Meat and offal. 13 day 13 days

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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:**

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

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**Available in:**

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

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**Package description:**

Packaging:Vial (glass), Package\_size:1 vial, Content:20 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:50 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:250 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Complete application (stand-alone) - Council Directive 81/851/EEC

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**Marketing authorisation holder:**

Zoetis Belgium

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**Marketing authorisation date:**

11/11/2003

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**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

11/11/2003

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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

Combined File of all Documents

English (PDF)

Published on: 25/04/2025

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ema-puar-v-077-var-x-0026-en.pdf

ema-puar-draxxin-v-077-var-x-0029-en.pdf

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ema-puar-draxxin-v-077-par-en.pdf

ema-puar-draxxin-v-077-var-ii-0034-en.pdf