

# Trilorale 10 mg/ml - Oral suspension

Authorised

- Trilostane

## Product identification

**Medicine name:**

Trilorale 10 mg/ml - Oral suspension

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**Active substance:**

Trilostane

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Trilostane

10.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral suspension

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

QH02CA01

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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:**

France

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**Package description:**

Packaging: Bottle with 1 ml and 5 ml PP syringe (HDPE), Package\_size: 1 bottle,  
Content: 30 ml

Packaging: Bottle with 1 ml and 5 ml PP syringe (HDPE), Package\_size: 1 bottle,  
Content: 90 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application - change in pharmaceutical form (Article 19(1)(a) of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

AXIENCE

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**Marketing authorisation date:**

6/05/2024

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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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:**

6/05/2024

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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)

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