# Trilorale 50 mg/ml - Oral suspension

Authorised

- Trilostane

## Product identification

**Medicine name:**
Trilorale 50 mg/ml - Oral suspension

**Active substance:**
Trilostane

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Trilostane
50.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Oral suspension

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QH02CA01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**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)

**Available in:**
France

**Package description:**
Packaging:Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size:1 bottle, Content:50 ml
Packaging:Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size:1 bottle, Content:100 ml
Packaging:Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size:1 bottle, Content:25 ml
Packaging:Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size:1 bottle, Content:10 ml
Packaging:Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size:1 bottle, Content:72 ml
Packaging:Bottle with 1 ml and 5 ml PP syringe (HDPE), Package_size:1 bottle, Content:36 ml

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**

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

**Marketing authorisation holder:**
AXIENCE

**Marketing authorisation date:**
6/05/2024

**Manufacturing sites for batch release:**
Lelypharma B.V.

**Responsible authority:**
European Commission

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
6/05/2024

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents

English (PDF)
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