

# Vetoryl 60 mg chewable tablets for dogs

Authorised

- Trilostane

## Product identification

**Medicine name:**

Vetoryl 60 mg chewable tablets for dogs

Vetoryl 60 mg purutabletti

**Active substance:**

Trilostane

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Trilostane

60.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Chewable tablet

**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:**

Finland

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**Package description:**

Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.

Cardboard box of 1 blister.

Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.

Cardboard box of 3 blisters.

Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.

Cardboard box of 5 blisters.

Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.

Cardboard box of 6 blisters.

Aluminium – Polyamide/Aluminium/PVC blister.Each blister contains 10 tablets.

Cardboard box of 10 blisters.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

10/10/2024

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

42716

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**Date of authorisation status change:**

10/10/2024

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0514/008

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Italy Latvia Lithuania Luxembourg  
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden 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

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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