

# Trilotab 10 mg chewable tablets for dogs

Authorised

- Trilostane

## Product identification

**Medicine name:**

Trilotab 10 mg chewable tablets for dogs

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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 Tablet

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**Pharmaceutical form:**

Chewable tablet

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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:**

Hungary

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**Package description:**

Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets. Cardboard box 100 tablets

Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets. Cardboard box of 30 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

CP-Pharma Handelsgesellschaft mbH

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**Marketing authorisation date:**

27/10/2023

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**Manufacturing sites for batch release:**

CP-Pharma Handelsgesellschaft mbH

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

4397/X/23 NÉBIH ÁTI

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**Date of authorisation status change:**

27/10/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0373/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Estonia Finland France Germany  
Hungary Ireland Italy Latvia Lithuania Poland Portugal Slovakia 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

Combined File of all Documents

PuAR Trilotab NL\_V\_0373\_001-005\_DC 2023-08.pdf