

# Trilotab 120 mg chewable tablets for

Authorised

- Trilostane

## Product identification

**Medicine name:**

Trilotab 120 mg chewable tablets for

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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

120.00 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:**

Lithuania

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**Available in:**

Lithuania

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**Package description:**

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

Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets. Cardboard box of 3 blister of 10 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:**

16/10/2023

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**Manufacturing sites for batch release:**

CP-Pharma Handelsgesellschaft mbH

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/23/2773/001-002

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**Date of authorisation status change:**

16/10/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0373/004

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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

English (PDF)

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