

# PROPALIN

Authorised

- Phenylpropanolamine hydrochloride

## Product identification

**Medicine name:**

PROPALIN

**Active substance:**

Phenylpropanolamine hydrochloride

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Phenylpropanolamine hydrochloride  
50.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Syrup

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

QG04BX

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovakia

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**Available in:**

Slovakia

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**Package description:**

Cardboard box with 1 bottle of 100 ml with a syringe of 1.5 ml

Cardboard box with 1 bottle of 30 ml with a syringe of 1.5 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetoquinol s.r.o.

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**Marketing authorisation date:**

2/06/2005

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**Manufacturing sites for batch release:**

Vetoquinol S.A.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/022/MR/05-S

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**Date of authorisation status change:**

2/06/2005

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0356/001

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**Concerned member states:**

Austria Belgium Cyprus Denmark Estonia Finland France Germany Greece  
Hungary Italy Latvia Lithuania Luxembourg Malta Poland Portugal Slovakia  
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

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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Package Leaflet