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

Authorisation status:
Valid

Authorised in:
Finland

Available in:
Finland

Package description:
Cardboard box with 1 bottle of 30 ml with a syringe of 1.5 ml
Cardboard box with 1 bottle of 100 ml with a syringe of 1.5 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
21/09/2003

Manufacturing sites for batch release:
Vetoquinol S.A.

Responsible authority:
Finnish Medicines Agency

Authorisation number:
17913

Date of authorisation status change:

21/09/2003

Reference member state:
Spain

Procedure number:
ES/V/0356/001

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)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/06/2025
Download

Package Leaflet

English (PDF)
Published on: 27/03/2026
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Combined File of all Documents