

# Urilin 40 mg/ml syrup for dogs

Authorised

- Phenylpropanolamine hydrochloride

## Product identification

**Medicine name:**

Urilin 40 mg/ml syrup for dogs

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**Active substance:**

Phenylpropanolamine hydrochloride

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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:**

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

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**Pharmaceutical form:**

Syrup

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

QG04BX91

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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:**

50 ml amber type III glass bottle containing 45 ml of syrup, with a low density polyethylene dropper and a polypropylene child resistant screw cap.

100 ml amber type III glass bottle containing 100 ml of syrup, with a low density polyethylene dropper and a polypropylene child resistant screw cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

21/09/2010

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**Manufacturing sites for batch release:**

Dales Pharmaceuticals Limited

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

2799/X/10 MgSzH ÁTI

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**Date of authorisation status change:**

21/09/2010

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0510/001

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**Concerned member states:**

Austria Czechia Denmark Germany Hungary Italy Portugal Slovakia Spain  
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