

Urilin 40 mg/ml syrup for dogs

Not authorised

- Phenylpropanolamine hydrochloride

Product identification

Medicine name:

Urilin 40 mg/ml syrup for dogs

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:

QG04BX91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Slovakia

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Dechra Regulatory B.V.

Marketing authorisation date:

5/08/2010

Manufacturing sites for batch release:

Dales Pharmaceuticals Limited

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/037/MR/10-S

Date of authorisation status change:

6/10/2023

Reference member state:

Ireland

Procedure number:

IE/V/0510/001

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics