Source URL: https://medicines.health.europa.eu/veterinary/en/60000050024

Urilin 40 mg/ml syrup for dogs

Not authorised

• Phenylpropanolamine hydrochloride

Product identification

Medicine name:

Urilin 40 mg/ml syrup for dogs Urilin, 40.0mg/ml, Sirup

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:

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Czechia

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:

3/11/2010

Manufacturing sites for batch release:

Dales Pharmaceuticals Limited

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/053/10-C

Date of authorisation status change: 17/07/2023
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
Summary of Product Characteristics
English (PDF) Published on: 3/05/2024 <u>Download</u>
Package Leaflet
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Labelling

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