

Urilin 40 mg/ml syrup for dogs

Not authorised

- Phenylpropanolamine hydrochloride

Product identification

Medicine name:

Urilin 40 mg/ml syrup for dogs

Urilin 40 mg/ml syrup

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:

Denmark

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:

14/06/2010

Manufacturing sites for batch release:

Dales Pharmaceuticals Limited

Responsible authority:

Danish Medicines Agency

Authorisation number:

46316

Date of authorisation status change:

24/07/2023

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)

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