Urilin 40 mg/ml syrup for dogs

Not authorised

- Phenylpropanolamine hydrochloride

Product identification

Medicine name:
Urilin 40 mg/ml syrup for dogs
URILIN SCIROPPO-REVOCATO

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:
Revoked

Authorised in:
Italy

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 Limited

Marketing authorisation date:
9/08/2010

Manufacturing sites for batch release:
Dales Pharmaceuticals Limited

Responsible authority:
Ministry Of Health

Authorisation number:
104251

Date of authorisation status change:
4/09/2010

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