Source URL: https://medicines.health.europa.eu/veterinary/en/700000102791

Uristop 40 mg/ml syrup for dogs

Authorised

• Phenylpropanolamine hydrochloride

Product identification

Medicine name:

Uristop 40 mg/ml syrup for dogs Urixine, 40mg/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

Withdrawal period by route of administration:

Oral use: Dog Anatomical therapeutic chemical veterinary (ATCvet) codes: OG04BX91 Legal status of supply: Veterinary medicinal product subject to veterinary prescription **Authorisation status:** Valid Authorised in: Czechia Package description: Box with 1 bottle of 100 ml and 1 syringe of 1,5 ml Box with 1 bottle of 50 ml and 1 syringe of 1,5 ml 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: Laboratorios Karizoo S.A. Marketing authorisation date: 17/07/2024

Manufacturing sites for batch release:

Fatro S.p.A.

Laboratorios Karizoo S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments
Authorisation number: 96/039/24-C
Date of authorisation status change: 17/07/2024
Reference member state: Spain
Procedure number: ES/V/0288/001
Concerned member states: Austria Belgium Czechia Estonia France Germany Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia United Kingdom (Northern Ireland)
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Package Leaflet
English (PDF) Published on: 5/06/2025

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