

# Uriphex 50 mg/ml, oral solution for dogs

Authorised

- Phenylpropanolamine hydrochloride

## Product identification

**Medicine name:**

Uriphex 50 mg/ml, oral solution for dogs

Uriphex 50 mg/ml Lösung zum Eingeben für Hunde

**Active substance:**

Phenylpropanolamine hydrochloride

**Target species:**

Dog (bitch)

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Phenylpropanolamine hydrochloride

50.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Oral solution

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QG04BX91

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Germany

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**Package description:**

HDPE bottle closed with white polypropylene child-resistant cap and LDPE syringe adaptor. A 1 mL HDPE/polypropylene graduated syringe is supplied with each bottle. Bottle of 100 mL.

HDPE bottle closed with white polypropylene child-resistant cap and LDPE syringe adaptor. A 1 mL HDPE/polypropylene graduated syringe is supplied with each bottle. Bottle of 60 mL.

HDPE bottle closed with white polypropylene child-resistant cap and LDPE syringe adaptor. A 1 mL HDPE/polypropylene graduated syringe is supplied with each bottle. Bottle of 30 mL.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

26/10/2023

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

V7009322.00.00

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**Date of authorisation status change:**

26/10/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0383/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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**Generic of:**

600000063880

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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