

# Telmitraxx 4 mg/ml oral solution for cats

Authorised

- Telmisartan

## Product identification

**Medicine name:**

Telmitraxx 4 mg/ml oral solution for cats

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**Active substance:**

Telmisartan

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**Target species:**

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Telmisartan

4.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral solution

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

QC09CA07

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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:**

Netherlands

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**Package description:**

HDPE bottle filled with 200 ml. Each bottle is closed with an LDPE plug-in adapter and a tamper-proof polypropylene (PP) closure. Pack size of one bottle and one measuring syringe (3 ml, LDPE barrel and piston, PS plunger).

HDPE bottle filled with 90 ml. Each bottle is closed with an LDPE plug-in adapter and a tamper-proof polypropylene (PP) closure. Pack size of one bottle and one measuring syringe (3 ml, LDPE barrel and piston, PS plunger).

HDPE bottle filled with 60 ml. Each bottle is closed with an LDPE plug-in adapter and a tamper-proof polypropylene (PP) closure. Pack size of one bottle and one measuring syringe (3 ml, LDPE barrel and piston, PS plunger).

HDPE bottle filled with 30 ml. Each bottle is closed with an LDPE plug-in adapter and a tamper-proof polypropylene (PP) closure. Pack size of one bottle and one measuring syringe (3 ml, LDPE barrel and piston, PS plunger).

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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:**

17/07/2023

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Alfasan Nederland B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 129763

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**Date of authorisation status change:**

24/08/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0386/001

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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:**

600000004056

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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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