

# Vominil 10 mg/ml solution for injection for dogs and cats

Authorised

- Maropitant citrate monohydrate

## Product identification

**Medicine name:**

Vominil 10 mg/ml Injektionslösung für Hunde und Katzen

Vominil 10 mg/ml solution for injection for dogs and cats

**Active substance:**

Maropitant citrate monohydrate

**Target species:**

Dog

Cat

**Route of administration:**

Intravenous use

Subcutaneous use

## Product details

**Active substance and strength:**

Maropitant citrate monohydrate

14.48 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Solution for injection

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

QA04AD90

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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:**

Ireland

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**Available in:**

Ireland

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**Package description:**

Amber glass vial type I (Ph. Eur.) with 10 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.

Amber glass vial type I (Ph. Eur.) with 25 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.

Amber glass vial type I (Ph. Eur.) with 50 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.

5 x Amber glass vials type I (Ph. Eur.) with 10 ml solution for injection, closed with a chlorobutyl rubber stopper, type I (Ph. Eur) and aluminium pull or flip off cap in a cardboard box.

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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:**

Vetviva Richter GmbH

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**Marketing authorisation date:**

29/09/2023

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**Manufacturing sites for batch release:**

Vetviva Richter GmbH

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA23462/018/001

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**Date of authorisation status change:**

29/09/2023

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0030/001

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**Concerned member states:**

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

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**Generic of:**

600000001521

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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