

# Diazedor 5 mg/ml Injektionslösung für Hunde und Katzen

Authorised

- Diazepam

## Product identification

**Medicine name:**

Diazedor 5 mg/ml Injektionslösung für Hunde und Katzen

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**Active substance:**

Diazepam

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**Target species:**

Dog

Cat

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Diazepam

5.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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

QN05BA01

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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:**

Sweden

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**Available in:**

Sweden

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**Package description:**

Colourless glass ampoules, type I, with 2 ml solution for injection, closed with a chlorobutyl rubber stopper and either an aluminium pull off cap or an aluminium/plastic flip off cap. Pack size: 10 x 2 ml in a cardboard box.

Colourless glass ampoules, type I, with 2 ml solution for injection, closed with a chlorobutyl rubber stopper and either an aluminium pull off cap or an aluminium/plastic flip off cap. Pack size: 5 x 2 ml in a cardboard box.

Colourless glass vials, type I, with 10 ml solution for injection, closed with a chlorobutyl rubber stopper and either an aluminium pull off cap or an aluminium/plastic flip off cap. Pack size: 1 x 10 ml 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:**

15/03/2018

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**Manufacturing sites for batch release:**

Vetviva Richter GmbH

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**Responsible authority:**

Swedish Medical Products Agency

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**Authorisation number:**

56075

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**Date of authorisation status change:**

15/03/2018

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0017/001

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**Concerned member states:**

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

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**Generic of:**

600000004401

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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

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Summary of Product Characteristics

English (PDF)

Published on: 13/03/2025

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### Labelling

This document does not exist in this language (English). You can find it in another language below.

### Combined File of all Documents

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