

# Zenalpha 0.5 mg + 10 mg - Solution for injection

Authorised

- Medetomidine hydrochloride
- Vatinoxan hydrochloride

## Product identification

**Medicine name:**

Zenalpha 0.5 mg + 10 mg - Solution for injection

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**Active substance:**

Medetomidine hydrochloride

Vatinoxan hydrochloride

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**Target species:**

Dog

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Medetomidine hydrochloride

Presentation\_strength:0.5 mg Reference:Hse Index:0

Vatinoxan hydrochloride

Presentation\_strength:10 mg Reference:Hse Index:1

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**Pharmaceutical form:**

Solution for injection

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

QN05CM99

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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:**

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

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**Available in:**

Austria , Belgium , Croatia , Denmark , Estonia , Finland , France , Germany , Ireland , Italy , Latvia , Lithuania , Netherlands , Norway , Poland , Portugal , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Vial (glass), Package\_size:10 vials, Content:10 ml

Packaging:Vial (glass), Package\_size:5 vials, Content:10 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:10 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetcare Oy

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**Marketing authorisation date:**

15/12/2021

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

Apotek Produktion & Laboratorier AB

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

15/12/2021

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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)

Published on: 22/12/2025

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