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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

Active substance:
Medetomidine hydrochloride

Vatinoxan hydrochloride

Target species:
Dog

Route of administration:
Intramuscular use

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

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

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)

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)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:10 ml

Packaging:Vial (glass), Package_size:5 vials, Content:10 ml

Packaging:Vial (glass), Package_size:10 vials, Content:10 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Vetcare Oy

Marketing authorisation date:
15/12/2021

Manufacturing sites for batch release:
Eurovet Animal Health B.V.
Apotek Produktion & Laboratorier AB

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
15/12/2021

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
Published on: 22/12/2025
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