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Varenzin 23.3 mg/ml - Oral suspension

Authorised

- Molidustat

Product identification

Medicine name:
Varenzin 23.3 mg/ml - Oral suspension

Active substance:
Molidustat

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Molidustat
23.30 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB03XA

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

Package description:
Packaging:Bottle (glass), syringe (PP), Package_size:1 bottle + 1 measuring syringe, Content:27 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - new active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Elanco GmbH

Marketing authorisation date:
22/01/2026

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
European Commission

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

Date of authorisation status change:
22/01/2026

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: 14/03/2026
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