

Numelvi 21.6 mg - Tablet

Authorised

- Atinvicitinib

Product identification

Medicine name:

Numelvi 21.6 mg - Tablet

Active substance:

Atinvicitinib

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Atinvicitinib

Presentation_strength:21.6 mg Reference:Hse Index:0

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD11AH93

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 , Germany

Package description:

Packaging:Cardboard box, Package_size:Cardboard box containing 1 blister strip with 30 tablets

Packaging:Cardboard box, Package_size:Cardboard box containing 3 blisters (30 each) contain 90 tablets in total

Packaging:Polyethylene bottle, Package_size:Polyethylene bottle containing 90 tablets

Packaging:Polyethylene bottle, Package_size:Polyethylene bottle containing 30 tablets

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:

Intervet International B.V.

Marketing authorisation date:

24/07/2025

Manufacturing sites for batch release:

Intervet Ges.m.b.H.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

24/07/2025

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

Documents

Combined File of all Documents

English (PDF)

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