

Lenivia 0.5 mg - Solution for injection

Authorised

- Izenivetmab

Product identification

Medicine name:

Lenivia 0.5 mg - Solution for injection

Active substance:

Izenivetmab

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Izenivetmab

Presentation_strength:0.45 - 0.55 mg Reference:In House Index:0

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02BG93

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:

Netherlands

Package description:

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

Packaging:Vial (Type I glass), Package_size:2 vials, Content:1 ml

Packaging:Vial (Type I glass), Package_size:6 vials, Content:1 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:

Zoetis Belgium

Marketing authorisation date:

21/11/2025

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

21/11/2025

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: 26/11/2025

[Download](#)