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:
Germany , 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)
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