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# Lenivia 2.0 mg - Solution for injection

Authorised

Izenivetmab

# **Product identification**

#### **Medicine name:**

Lenivia 2.0 mg - Solution for injection

#### **Active substance:**

Izenivetmab

#### **Target species:**

Dog

#### **Route of administration:**

Subcutaneous use

# **Product details**

### **Active substance and strength:**

Izenivetmab

Presentation\_strength:1.8 - 2.2 mg Reference:In House Index:0

#### **Pharmaceutical form:**

Solution for injection

# Withdrawal period by route of administration: Subcutaneous use:

Dog

#### 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)

#### Package description:

Packaging:Vial (Type I glass), Package\_size:1 vial, Content:1 ml Packaging:Vial (Type I glass), Package\_size:6 vials, Content:1 ml Packaging:Vial (Type I glass), Package\_size:2 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 <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Combined File of all Documents

English (PDF)

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