

# Librela 10 mg - Solution for injection

Authorised

- Bedinvetmab

## Product identification

**Medicine name:**

Librela 10 mg - Solution for injection

**Active substance:**

Bedinvetmab

**Target species:**

Dog

**Route of administration:**

Subcutaneous use

## Product details

**Active substance and strength:**

Bedinvetmab

Presentation\_strength:9 - 11 mg Reference:Hse Index:0

**Pharmaceutical form:**

Solution for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Dog**

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN02BG91

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Available in:**

Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Vial (glass), Package\_size:6 vials, Content:1 ml

Packaging:Vial (glass), Package\_size:2 vials, Content:1 ml

Packaging:Vial (glass), Package\_size:1 vial, Content:1 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Belgium SA

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**Marketing authorisation date:**

10/11/2020

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**Manufacturing sites for batch release:**

Zoetis Belgium

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

10/11/2020

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 12/02/2025

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