

# DALMARELIN 25 micrograms/ml solution for injection for cattle and rabbits

Authorised

- Lecirelin

## Product identification

**Medicine name:**

DALMARELIN 25 micrograms/ml solution for injection for cattle and rabbits

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**Active substance:**

Lecirelin

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**Target species:**

Rabbit

Cattle

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Lecirelin

25.00 microgram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Rabbit**

- Meat and offal. 0 day

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**Cattle**

- Meat and offal. 0 day

- Milk. 0 day

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

QH01CA92

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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:**

France

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**Available in:**

France

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**Package description:**

100 ml High Density Polyethylene (HDPE) collapsible container closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

20 ml type II neutral colourless glass vials, closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

5 x 10 ml type I neutral colourless glass vials, closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

10 ml type I neutral colourless glass vials, closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

10 x 4 ml type I neutral colourless glass vials, closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

4 ml type I neutral colourless glass vials, closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

50 ml type II neutral colourless glass vials, closed with a type I rubber stopper and an aluminium overseal, in a cardboard box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

23/01/2004

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/2552317 9/2004

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**Date of authorisation status change:**

23/01/2009

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**Reference member state:**

Italy

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**Procedure number:**

IT/V/0112/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland France  
Germany Hungary Iceland Ireland Luxembourg Malta Netherlands Norway  
Poland Portugal Romania Slovakia Slovenia Spain Sweden

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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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents