

# 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:**

Ireland

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**Available in:**

Ireland

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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:**

19/03/2004

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10836/002/001

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**Date of authorisation status change:**

19/03/2004

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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

Combined File of all Documents