

# DALMARELIN 25 MCG/ML, ενέσιμο διάλυμα για αγελάδες, φοράδες και κουνέλια

Authorised

- Lecirelin

## Product identification

**Medicine name:**

DALMARELIN 25 MCG/ML, ενέσιμο διάλυμα για αγελάδες, φοράδες και κουνέλια

**Active substance:**

Lecirelin

**Target species:**

Cattle (cow)

Rabbit

Horse (mare)

**Route of administration:**

Intramuscular and intravenous use

## Product details

**Active substance and strength:**

Lecirelin

25.00 microgram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Solution for injection

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

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**Cattle (cow)**

- Meat and offal. 0 day

- Milk. 0 hour

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**Rabbit**

- Meat and offal. 0 day

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**Horse (mare)**

- Meat and offal. 0 day

- Milk. 0 hour

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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:**

Greece

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**Available in:**

Greece

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**Package description:**

Available only in Greek

Available only in Greek

Available only in Greek

Available only in [Greek](#)

Available only in [Greek](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Complete application (stand-alone)

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

9/03/2000

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

9681/10-03-2000/K-0126901

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**Date of authorisation status change:**

18/06/2024

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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.