

# Leventa 1 mg/ml oral solution for dogs

Authorised

- Levothyroxine sodium

## Product identification

**Medicine name:**

Leventa 1 mg/ml oral solution for dogs

Leventa 1 mg/ml Roztwór doustny

**Active substance:**

Levothyroxine sodium

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Levothyroxine sodium

1.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Oral solution

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

QH03AA01

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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:**

Poland

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**Package description:**

30 mL amber glass bottle with a transparent LDPE insert and with a white HDPE child-resistant cap with tamper-proofclosure in a printed carton.A 1 ml oral syringe graduated in 0.05 ml increments is supplied with the product.Pack size: 1 x 30 ml  
30 mL amber glass bottle with a transparent LDPE insert and with a white HDPE child-resistant cap with tamper-proofclosure in a printed carton.A 1 ml oral syringe graduated in 0.05 ml increments is supplied with the product.Pack size: 6 x 30 ml  
30 mL amber glass bottle with a transparent LDPE insert and with a white HDPE child-resistant cap with tamper-proofclosure in a printed carton.A 1 ml oral syringe graduated in 0.05 ml increments is supplied with the product.Pack size: 12 x 30 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

13/08/2007

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**Manufacturing sites for batch release:**

Intervet Productions S.A.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1765

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**Date of authorisation status change:**

13/08/2007

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0182/001

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**Concerned member states:**

Austria Belgium Czechia France Germany Greece Hungary Italy  
Luxembourg Netherlands Poland Portugal Spain Sweden  
United Kingdom (Northern Ireland)

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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

Package Leaflet

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

Summary of Product Characteristics

English (PDF)

Published on: 1/06/2025

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Combined File of all Documents