# Leventa 1 mg/ml oral solution for dogs

Not authorised

- Levothyroxine sodium

## Product identification

**Medicine name:**
Leventa 1 mg/ml oral solution for dogs
Leventa, 1mg/ml, Perorální roztok

**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

**Legal status of supply:**
Medicinal product subject to medical prescription

**Authorisation status:**
Surrendered

**Authorised in:**
Czechia

**Package description:**
30 mL amber glass bottle with a transparent LDPE insert and with a white HDPE child-resistant cap with tamper-proof closure 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-proof closure 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-proof closure 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

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Intervet International B.V.

**Marketing authorisation date:**
25/06/2007

**Manufacturing sites for batch release:**
Intervet Productions S.A.

**Responsible authority:**
Institute For State Control Of Veterinary Biologicals And Medicaments

**Authorisation number:**
96/026/07-C

**Date of authorisation status change:**
28/08/2023

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0182/001

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 1/06/2025
Download

Package Leaflet

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Labelling

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

Combined File of all Documents