

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 Lösung zum Eingeben für Hunde

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

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

-

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH03AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

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

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

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 Ges.m.b.H.

Marketing authorisation date:

4/05/2007

Manufacturing sites for batch release:

Intervet Productions S.A.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-00700

Date of authorisation status change:

4/05/2007

Reference member state:

Ireland

Procedure number:

IE/V/0182/001

Concerned member states:

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

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

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Package Leaflet

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Labelling

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