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 1 mg/ml oplossing voor oraal gebruik bij honden

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:
Surrendered

Authorised in:
Netherlands

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 Nederland B.V.

Marketing authorisation date:
6/11/2007

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 10408

Date of authorisation status change:
29/08/2023

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

Combined File of all Documents

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

Summary of Product Characteristics
