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 belsőleges oldat kutyáknak

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:

Hungary

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

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:
12/04/2007

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

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
2179/X/07 MgSzH ÁTI

Date of authorisation status change:
12/04/2007

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

Combined File of all Documents