# Nobivac Lepto

Authorised

- Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain Ic-02-001, Inactivated
- Leptospira interrogans, serogroup Canicola, serovar Portland-vere, strain Ca-12-000, Inactivated

## Product identification

**Medicine name:**
NOBIVAC LEPTO ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ
Nobivac Lepto

**Active substance:**
Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain Ic-02-001, Inactivated

Leptospira interrogans, serogroup Canicola, serovar Portland-vere, strain Ca-12-000, Inactivated

**Target species:**
Dog

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain Ic-02-001, Inactivated

699.00 unit(s) / 1.00 Dose

Leptospira interrogans, serogroup Canicola, serovar Portland-vere, strain Ca-12-000, Inactivated
990.00 unit(s) / 1.00 Dose

**Pharmaceutical form:**
Suspension for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Dog**

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI07AB01

**Legal status of supply:**
This information is not available for this product.

**Authorisation status:**
Valid

**Authorised in:**
Greece

**Package description:**
Plasic box containing 50 type I glass vials of 1 ml (1 dose) closed with a halogenobutyl rubber stopper and sealed with a coded aluminium cap
Plasic box containing 10 type I glass vials of 1 ml (1 dose) closed with a halogenobutyl rubber stopper and sealed with a coded aluminium cap
Cardboard box containing 50 type I glass vials of 1 ml (1 dose) closed with a halogenobutyl rubber stopper and sealed with a coded aluminium cap
Cardboard box containing 10 type I glass vials of 1 ml (1 dose) closed with a halogenobutyl rubber stopper and sealed with a coded aluminium cap

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Legal basis not covered by Directive 2001/82/EC

**Marketing authorisation holder:**
Intervet Hellas A.E.

**Marketing authorisation date:**
28/07/2003

**Manufacturing sites for batch release:**
INTERVET INTERNATIONAL B.V.

**Responsible authority:**
National Organization For Medicines

**Authorisation number:**
56940/08-09-2008/K-0148301

**Date of authorisation status change:**
8/09/2008

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0108/001

**Concerned member states:**
Belgium Denmark Greece Luxembourg Portugal Spain

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

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