

Nobivac Lepto suspension for injection for dogs

Authorised

- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
- Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

Product identification

Medicine name:

Nobivac Lepto suspension for injection for dogs

Active substance:

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated

990.00 unit(s) / 1.00 millilitre(s)

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated
699.00 unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

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

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

Plastic 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

Plastic 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

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 M.A.E.

Marketing authorisation date:

27/07/2003

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:

7/09/2008

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Combined File of all Documents