Nobivac Lepto

Not 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
Nobivac Lepto Suspensie voor injectie
Nobivac Lepto Suspension injectable
Nobivac Lepto Injektionssuspension

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:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
Belgium

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

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Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
12/01/2004

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V259025

Date of authorisation status change:
22/08/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0108/001

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.
