Nobivac RL suspensija injekcijām suņiem

Authorised

- Rabies virus, strain Pasteur RIV, Inactivated
- Leptospira interrogans, serogroup Canicola, serovar Portland-vere, strain Ca-12-000, Inactivated
- Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain 820K, Inactivated

Product identification

Medicine name:
Nobivac RL suspensija injekcijām suņiem

Active substance:
Rabies virus, strain Pasteur RIV, Inactivated

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

Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain 820K, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Rabies virus, strain Pasteur RIV, Inactivated
3.00 international unit(s) / 1.00 unit(s)

Leptospira interrogans, serogroup Canicola, serovar Portland-vere, strain Ca-12-000, Inactivated
4.00 Hamster protective Dose 80 % (Ph. Eur. Monograph) / 1.00 unit(s)

Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain 820K, Inactivated
4.00 Hamster protective Dose 80 % (Ph. Eur. Monograph) / 1.00 unit(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Latvia

Package description:

Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
22/03/1996

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

Responsible authority:
PVD

Authorisation number:
V/NRP/96/0379

Date of authorisation status change:
24/03/1996

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.

Labelling

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