

Nobivac RL, injekciné suspensija šunims

Authorised

- Rabies virus, strain Pasteur RIV, Inactivated
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
- Leptospira interrogans, serovar Copenhageni, strain 820K, Inactivated

Product identification

Medicine name:

Nobivac RL, injekciné suspensija šunims

Active substance:

Rabies virus, strain Pasteur RIV, Inactivated

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

Leptospira interrogans, 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 millilitre(s)

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

40.00 Hamster protective Dose 80% / 1.00 millilitre(s)

Leptospira interrogans, serovar Copenhageni, strain 820K, Inactivated

40.00 Hamster protective Dose 80% / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Available in:

Lithuania

Package description:

Available only in [Lithuanian](#)

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:

20/02/2000

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/00/1069/001

Date of authorisation status change:

21/09/2010

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

Documents

Combined File of all Documents

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