Nobivac RL

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

Authorised

Product identification

Medicine name:
Nobivac RL
Active substance:

- Rabies virus, strain Pasteur RIV, Inactivated
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, 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
  dose
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
  40.00
  Hamster protective Dose 80%
  /
  1.00
  dose

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:

- Estonia

Package description:

- Available only in Estonian

Additional information

Entitlement type:

- Marketing Authorisation

Legal basis of product authorisation:

- Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

- Intervet International B.V.

Marketing authorisation date:

- 13/10/2005

Manufacturing sites for batch release:

- Intervet International B.V.

Responsible authority:

- State Agency Of Medicines

Authorisation number:

- 1339

Date of authorisation status change:

- 13/10/2005

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

Documents

Product information

Summary of Product Characteristics

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