

NOBIVAC RL ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ

Authorised

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

Product identification

Medicine name:

NOBIVAC RL ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ

Active substance:

Rabies virus, strain Pasteur RIV, Inactivated

Leptospira interrogans, serovar Copenhageni, strain 820K, 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 millilitre(s)

Leptospira interrogans, serovar Copenhageni, strain 820K, Inactivated

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

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

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

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Dog

- Not applicable. no withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Available only in [Greek](#)

Available only in [Greek](#)

Available only in [Greek](#)

Available only in [Greek](#)

Available only in [Greek](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Intervet Hellas M.A.E.

Marketing authorisation date:

13/01/1993

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

National Organization For Medicines

Authorisation number:

27950/97/K2520/27-1-1998/K-0076101

Date of authorisation status change:

15/07/2021

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