

Nobivac L + DHPPi lyofilisaat en suspensie voor suspensie voor injectie voor honden

Authorised

- Canine distemper virus, strain Onderstepoort, Live
- Canine parainfluenza virus, strain Cornell, Live
- Canine parvovirus, strain 154, Live
- Canine adenovirus 2, strain Manhattan LPV3, Live
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
- Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

Product identification

Medicine name:

Nobivac L + DHPPi lyofilisaat en suspensie voor suspensie voor injectie voor honden

Active substance:

Canine distemper virus, strain Onderstepoort, Live

Canine parainfluenza virus, strain Cornell, Live

Canine parvovirus, strain 154, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

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

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine distemper virus, strain Onderstepoort, Live

4.00 log₁₀ 50% cell culture infectious dose / 1.00 millilitre(s)

Canine parainfluenza virus, strain Cornell, Live

5.50 log₁₀ 50% cell culture infectious dose / 1.00 millilitre(s)

Canine parvovirus, strain 154, Live

7.00 log₁₀ 50% cell culture infectious dose / 1.00 millilitre(s)

Canine adenovirus 2, strain Manhattan LPV3, Live

4.00 log₁₀ 50% cell culture infectious dose / 1.00 millilitre(s)

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

1300.00 enzyme-linked immunosorbent assay unit/millilitre / 1.00 millilitre(s)

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

750.00 enzyme-linked immunosorbent assay unit/millilitre / 1.00 millilitre(s)

Pharmaceutical form:

Lyophilisate for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AI02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

Available only in Dutch

Available only in [Dutch](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet Nederland B.V.

Marketing authorisation date:

9/10/2002

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 9483

Date of authorisation status change:

28/03/2013

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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