

Vanguard Plus 5/L EU, Lyofilizát a rozpouštědlo pro injekční suspenzi

Authorised

- Canine distemper virus, strain N-CDV, Live
- Canine adenovirus 2, strain Manhattan, Live
- Canine parainfluenza virus, strain NL-CPI-5, Live
- Canine parvovirus, strain NL-35-D, Live
- Leptospira interrogans, serogroup Canicola, serovar Canicola, strain C51, Inactivated
- Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Product identification

Medicine name:

Vanguard Plus 5/L EU, Lyofilizát a rozpouštědlo pro injekční suspenzi

Active substance:

Canine distemper virus, strain N-CDV, Live

Canine adenovirus 2, strain Manhattan, Live

Canine parainfluenza virus, strain NL-CPI-5, Live

Canine parvovirus, strain NL-35-D, Live

Leptospira interrogans, serogroup Canicola, serovar Canicola, strain C51, Inactivated

Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine distemper virus, strain N-CDV, Live

5.50 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, strain Manhattan, Live

5.20 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine parainfluenza virus, strain NL-CPI-5, Live

8.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine parvovirus, strain NL-35-D, Live

8.50 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Leptospira interrogans, serogroup Canicola, serovar Canicola, strain C51, Inactivated

740.00 relative potency / 1.00 Dose

Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar

Icterohaemorrhagiae, strain NADL 11403, Inactivated

915.00 relative potency / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:**Subcutaneous use:**

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Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AI02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

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:

Zoetis Ceska Republika s.r.o.

Marketing authorisation date:

17/11/1992

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/668/92-S/C/02

Date of authorisation status change:

1/10/2009

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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