Vanguard Plus 5/L EU lyofilizát a tekutá zložka na injekčnú suspenziu pre psy

Authorised

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

Product identification

Medicine name:
Vanguard Plus 5/L EU lyofilizát a tekutá zložka na injekčnú suspenziu pre psy

Active substance:
Canine adenovirus 2, strain Manhattan, Live

Canine distemper virus, strain N-CDV, Live

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

Canine parvovirus, strain NL-35-D, Live

Leptospira interrogans, serovar Canicola, strain C51, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Target species:

Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine adenovirus 2, strain Manhattan, Live
3.20 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine distemper virus, strain N-CDV, Live
3.00 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parainfluenza virus, strain NL-CPI-5, Live
6.00 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parvovirus, strain NL-35-D, Live
7.00 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Leptospira interrogans, serovar Canicola, strain C51, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Pharmaceutical form:
Lyophilisate and suspension for suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Dog**
  - All relevant tissues. 0 day not applicable

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:
Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak

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

Date of authorisation status change:
17/11/1992

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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