VANGUARD PLUS 7

Authorised

- Canine parainfluenza virus, strain NL-CPI-5, Live
- Canine distemper virus, strain N-CDV, Live
- Canine adenovirus 2, strain Manhattan, 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 7

Active substance:
Canine parainfluenza virus, strain NL-CPI-5, Live

Canine distemper virus, strain N-CDV, Live

Canine adenovirus 2, strain Manhattan, 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
Dog

Route of administration:

Subcutaneous use
Subcutaneous use

Product details

Active substance and strength:
Canine parainfluenza virus, strain NL-CPI-5, Live
6.00 log10 50% cell culture infectious dose / 1.00 millilitre(s)

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

Canine adenovirus 2, strain Manhattan, Live
3.20 log10 50% cell culture infectious dose / 1.00 millilitre(s)

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

Leptospira interrogans, serovar Canicola, strain C51, Inactivated
40.00 Protective Dose / 1.00 millilitre(s)

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated
40.00 Protective Dose / 1.00 millilitre(s)

Pharmaceutical form:
Lyophilisate and solvent for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AJ02

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

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:

Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
19/06/2007

Manufacturing sites for batch release:
Zoetis Belgium SA

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
140019

Date of authorisation status change:
26/10/2023

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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