Vanguard plus 7 liofilizat in vehikel za raztopino za injiciranje za pse

Authorised

- LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED
- Leptospira interrogans, serovar Canicola, Inactivated
- Canine parvovirus, strain NL-35-D, Live
- Canine parainfluenza virus, strain NL-CPI-5, Live
- Canine adenovirus 2, strain Manhattan, Live
- Canine distemper virus, strain N-CDV, Live

Product identification

Medicine name:
Vanguard plus 7 liofilizat in vehikel za raztopino za injiciranje za pse

Active substance:
LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED
Leptospira interrogans, serovar Canicola, Inactivated
Canine parvovirus, strain NL-35-D, Live
Canine parainfluenza virus, strain NL-CPI-5, Live
Canine adenovirus 2, strain Manhattan, Live
Canine distemper virus, strain N-CDV, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED
40.00 Protective Dose / 1.00 millilitre(s)

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

Canine parvovirus, strain NL-35-D, Live
10000000.00 50% cell culture infectious dose / 1.00 millilitre(s)

Canine parainfluenza virus, strain NL-CPI-5, Live
1000000.00 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, strain Manhattan, Live
1584.00 50% cell culture infectious dose / 1.00 Dose

Canine distemper virus, strain N-CDV, Live
1000.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for solution 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:
Slovenia

Available in:
Slovenia

Package description:

Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
10/11/2021

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0347/001

Date of authorisation status change:
28/09/2004

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.