VANGUARD 7

Not authorised

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

Product identification

Medicine name:
VANGUARD 7

Active substance:
Canine adenovirus 2, strain Manhattan, Live

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

Canine distemper virus, strain N-CDV, Live

Canine parvovirus, strain NL-35-D, Live

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Leptospira interrogans, serovar Canicola, strain C51, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Canine adenovirus 2, strain Manhattan, Live
1584.89 50% tissue culture infectious dose / 1.00 unit(s)

Canine parainfluenza virus, strain NL-CPI-5, Live
5011870.00 50% tissue culture infectious dose / 1.00 unit(s)

Canine distemper virus, strain N-CDV, Live
1000.00 50% tissue culture infectious dose / 1.00 unit(s)

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

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated
463.00 relative unit(s) / 1.00 unit(s)

Leptospira interrogans, serovar Canicola, strain C51, Inactivated
420.00 relative unit(s) / 1.00 unit(s)

Pharmaceutical form:

Lyophilisate and suspension for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AI02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

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Authorisation status:

Surrendered

Authorised in:

France

Package description:

Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis France

Marketing authorisation date:
15/09/1995

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6862556 2/1995

Date of authorisation status change:
21/10/2022

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.