

# VANGUARD PLUS 7, liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań dla psów

Authorised

- Canine adenovirus 2, Live
- Canine parainfluenza virus, Live
- Canine parvovirus, Live
- Leptospira interrogans, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Canine distemper virus, Live

## Product identification

### **Medicine name:**

VANGUARD PLUS 7, liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań dla psów

### **Active substance:**

Canine adenovirus 2, Live

Canine parainfluenza virus, Live

Canine parvovirus, Live

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine distemper virus, Live

**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Canine adenovirus 2, Live

1584.89 50% cell culture infectious dose / 1.00 millilitre(s)

Canine parainfluenza virus, Live

1000000.00 50% cell culture infectious dose / 1.00 millilitre(s)

Canine parvovirus, Live

10000000.00 50% cell culture infectious dose / 1.00 millilitre(s)

Leptospira interrogans, serovar Canicola, Inactivated

740.00 relative unit(s) / 1.00 millilitre(s)

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

915.00 relative unit(s) / 1.00 millilitre(s)

Canine distemper virus, Live

1000.00 50% cell culture infectious dose / 1.00 millilitre(s)

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**Pharmaceutical form:**

Lyophilisate and solvent for suspension for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI07AI02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**Package description:**

Available only in Polish

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Polska Sp. z o.o.

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**Marketing authorisation date:**

30/04/2004

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**Manufacturing sites for batch release:**

Zoetis Belgium SA

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1461

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**Date of authorisation status change:**

30/04/2004

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

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### Package Leaflet

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