

# Vanguard Plus 5 vakcina A.U.V.

Authorised

- 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 5 vakcina A.U.V.

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**Active substance:**

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

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**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Canine parvovirus, strain NL-35-D, Live

7.00 log10 50% cell culture infectious dose / 1.00 unit(s)/dose

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

6.00 log10 50% cell culture infectious dose / 1.00 unit(s)/dose

Canine adenovirus 2, strain Manhattan, Live

3.20 log10 50% cell culture infectious dose / 1.00 unit(s)/dose

Canine distemper virus, strain N-CDV, Live

3.00 log10 50% cell culture infectious dose / 1.00 unit(s)/dose

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**Pharmaceutical form:**

Powder and solvent for suspension for injection

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

QI07AD04

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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:**

Hungary

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**Available in:**

Hungary

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**Package description:**

Available only in Hungarian

Available only in Hungarian

Available only in Hungarian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Hungary Kft.

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**Marketing authorisation date:**

30/04/2004

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**Manufacturing sites for batch release:**

Zoetis Belgium

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

This information is not available for this product.

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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  
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