

Vanguard Plus Puppy lyofilizát a tekutá zložka na injekčnú suspenziu pre psy

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 Puppy lyofilizát a tekutá zložka na injekčnú suspenziu pre psy

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

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine parvovirus, strain NL-35-D, Live

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

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

6.00 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 distemper virus, strain N-CDV, Live

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

Pharmaceutical form:

Lyophilisate and suspension for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Available in:

Slovakia

Package description:

Available only in Slovak

Available only in Slovak

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Zoetis Ceska Republika s.r.o.

Marketing authorisation date:

30/04/2004

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/083/04-S

Date of authorisation status change:

30/04/2004

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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