

CANIGEN DHA2PPiLR, Lyofilizát a rozpouštědlo pro injekční suspenzi

Authorised

- Canine distemper virus, Live
- Canine adenovirus 2, Live
- Canine parvovirus, Live
- Canine parainfluenza virus, Live
- Rabies virus, strain VP12, Inactivated
- Leptospira interrogans, serovar Canicola, strain 601903, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated

Product identification

Medicine name:

CANIGEN DHA2PPiLR, Lyofilizát a rozpouštědlo pro injekční suspenzi

Active substance:

Canine distemper virus, Live

Canine adenovirus 2, Live

Canine parvovirus, Live

Canine parainfluenza virus, Live

Rabies virus, strain VP12, Inactivated

Leptospira interrogans, serovar Canicola, strain 601903, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine distemper virus, Live

5.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live

6.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, Live

7.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Canine parainfluenza virus, Live

7.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Rabies virus, strain VP12, Inactivated

1.00 international unit(s) / 1.00 millilitre(s)

Leptospira interrogans, serovar Canicola, strain 601903, Inactivated

40.00 Protective Dose / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated

40.00 Protective Dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AJ06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

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:

Virbac

Marketing authorisation date:

9/09/1994

Manufacturing sites for batch release:

Virbac

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/962/94-C

Date of authorisation status change:

9/09/2010

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.