

CANVAC DHPPiL+R, liofilizatas ir skiediklis injekcinei suspensijai

Authorised

- Rabies virus, Inactivated
- LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED
- Canine parainfluenza virus, Live
- Canine parvovirus, Live
- Canine adenovirus 2, Live
- Canine distemper virus, Live

Product identification

Medicine name:

CANVAC DHPPiL+R, liofilizatas ir skiediklis injekcinei suspensijai

Active substance:

Rabies virus, Inactivated

LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED

Canine parainfluenza virus, Live

Canine parvovirus, Live

Canine adenovirus 2, Live

Canine distemper virus, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabies virus, Inactivated

2.00 international unit(s) / 1.00 Dose

LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED

32.00 antibody unit(s) / 1.00 Dose

Canine parainfluenza virus, Live

1000.00 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, Live

128.00 haemagglutinating units / 1.00 Dose

Canine adenovirus 2, Live

3162.28 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, Live

501.19 50% Embryo Infective 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:

Lithuania

Package description:

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Dyntec spol. s r.o.

Marketing authorisation date:

23/06/2009

Manufacturing sites for batch release:

Dyntec spol. s r.o.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

6/08/2014

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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