

CANIGEN DHA2PPi /LR, Lyophilisate and solvent for suspension for injection

Authorised

- Rabies virus, strain VP12, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated
- Leptospira interrogans, serovar Canicola, strain 601903, Inactivated
- Canine parainfluenza virus, strain Manhattan, Live
- Canine parvovirus, strain Cornell 780916, Live
- Canine adenovirus 2, strain Manhattan, Live
- Canine distemper virus, strain Lederle, Live

Product identification

Medicine name:

CANIGEN DHA2PPi /LR, Lyophilisate and solvent for suspension for injection

Active substance:

Rabies virus, strain VP12, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated

Leptospira interrogans, serovar Canicola, strain 601903, Inactivated

Canine parainfluenza virus, strain Manhattan, Live

Canine parvovirus, strain Cornell 780916, Live

Canine adenovirus 2, strain Manhattan, Live

Canine distemper virus, strain Lederle, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabies virus, strain VP12, Inactivated

1.00 international unit(s) / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated

8.33 80% Protective Dose / 1.00 Dose

Leptospira interrogans, serovar Canicola, strain 601903, Inactivated

8.33 80% Protective Dose / 1.00 Dose

Canine parainfluenza virus, strain Manhattan, Live

5.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine parvovirus, strain Cornell 780916, Live

5.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, strain Manhattan, Live

4.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine distemper virus, strain Lederle, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AI01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in Bulgarian

Available only in Bulgarian

Available only in Bulgarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Virbac

Marketing authorisation date:

8/03/1999

Manufacturing sites for batch release:

Virbac

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2296

Date of authorisation status change:

8/03/1999

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.