

VANGUARD CPV-L.

Authorised

- Leptospira interrogans, serovar Canicola, strain C51, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated
- Canine parvovirus, strain NL-35-D, Live

Product identification

Medicine name:

VANGUARD CPV-L.

Active substance:

Leptospira interrogans, serovar Canicola, strain C51, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Canine parvovirus, strain NL-35-D, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Canicola, strain C51, Inactivated

40.00 unit(s) / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

40.00 unit(s) / 1.00 Dose

Canine parvovirus, strain NL-35-D, Live

316228000.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AI05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

The vaccine is filled in 1 dose vials glass type I (Ph.Eur.). Pack contains 100vials of 1 ml liquid VanguardCPV-L.

The vaccine is filled in 1 dose vials glass type I (Ph.Eur.). Pack contains 25vials of 1 ml liquid VanguardCPV-L.

The vaccine is filled in 1 dose vials glass type I (Ph.Eur.). Pack contains 10vials of 1 ml liquid VanguardCPV-L.

The vaccine is filled in 1 dose vials glass type I (Ph.Eur.). Pack contains 1vial of 1 ml liquid VanguardCPV-L.

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 Belgium S.A.

Marketing authorisation date:

12/09/2014

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10387/084/001

Date of authorisation status change:

12/09/2014

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

Documents

Summary of Product Characteristics