

TETRAVALENT L-CHP

Authorised

- Canine parvovirus, strain CPV-N-1988, Live
- Canine adenovirus 2, Live
- Canine distemper virus, strain CDV/WHO 134, Live
- Leptospira interrogans, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Canine parvovirus, strain CPV-N-1988, Live
- Canine adenovirus 2, Live
- Canine distemper virus, strain CDV/WHO 134, Live

Product identification

Medicine name:

TETRAVALENT L-CHP

Active substance:

Canine parvovirus, strain CPV-N-1988, Live

Canine adenovirus 2, Live

Canine distemper virus, strain CDV/WHO 134, Live

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine parvovirus, strain CPV-N-1988, Live

Canine adenovirus 2, Live

Canine distemper virus, strain CDV/WHO 134, Live

Target species:

Dog

Dog

Route of administration:

Subcutaneous use

Subcutaneous use

Product details

Active substance and strength:

Canine parvovirus, strain CPV-N-1988, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine distemper virus, strain CDV/WHO 134, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Leptospira interrogans, serovar Canicola, Inactivated

100.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

100.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Canine parvovirus, strain CPV-N-1988, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Canine distemper virus, strain CDV/WHO 134, Live

3.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AI03

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Romvac Company S.A.

Marketing authorisation date:

15/03/2007

Manufacturing sites for batch release:

Romvac Company S.A.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

120233

Date of authorisation status change:

7/11/2006

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

Documents

Summary of Product Characteristics

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