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Versican Plus DHPPi/L4 (--) - Lyophilisate and suspension for suspension for injection

Authorised

- Canine distemper virus, strain CDV Bio 11/A, Live
- Canine adenovirus 2, strain CAV-2-Bio 13, Live
- Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live
- Canine parainfluenza virus, strain CPiV-2-Bio 15, Live
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1089, Inactivated
- Leptospira interrogans, serovar Bratislava, strain MSLB 1088, Inactivated
- Leptospira interrogans, serovar Canicola, strain MSLB 1090, Inactivated
- Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1091, Inactivated

Product identification

Medicine name:
Versican Plus DHPPi/L4 (--) - Lyophilisate and suspension for suspension for injection

Active substance:
Canine distemper virus, strain CDV Bio 11/A, Live

Canine adenovirus 2, strain CAV-2-Bio 13, Live

Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live

Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1089, Inactivated

Leptospira interrogans, serovar Bratislava, strain MSLB 1088, Inactivated

Leptospira interrogans, serovar Canicola, strain MSLB 1090, Inactivated

Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1091, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, strain CDV Bio 11/A, Live

Presentation_strength:$10^{3,1}$ - $10^{5,1}$ $TCID_{50}$ Reference:Ph Eur 0448 Comments:Lyophilisate Index:0

Canine adenovirus 2, strain CAV-2-Bio 13, Live

Presentation_strength:$10^{3,6}$ - $10^{5,3}$ $TCID_{50}$ Reference:Ph Eur 1951 Comments:Lyophilisate Index:1

Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live

Presentation_strength:$10^{4,3}$ - $10^{6,6}$ $TCID_{50}$ Reference:Ph Eur 0964 Comments:Lyophilisate Index:2

Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Presentation_strength:$10^{3,1}$ - $10^{5,1}$ $TCID_{50}$1 Reference:Ph Eur 1955 Comments:Lyophilisate Index:3

Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1089, Inactivated

Presentation_strength:≥1:51 ALR Reference:Ph Eur 0447 Comments:Suspension Index:4

Leptospira interrogans, serovar Bratislava, strain MSLB 1088, Inactivated

Presentation_strength:≥1:51 ALR Reference:Ph Eur 0447 Comments:Suspension Index:5

Leptospira interrogans, serovar Canicola, strain MSLB 1090, Inactivated

Presentation_strength:≥1:51 ALR Reference:Ph Eur 0447 Comments:Suspension Index:6

Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1091, Inactivated Presentation_strength:≥1:40 ALR Reference:Ph Eur 0447 Comments:Suspension Index:7

Pharmaceutical form:
Lyophilisate and suspension 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:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:
Austria , Belgium , Bulgaria , Cyprus , Czechia , Denmark , Estonia , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Poland , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

Package description:
Packaging:Lyophilisate: vial (glass); suspension: vial (glass), Package_size:Lyophilisate: 50 vials + suspension: 50 vials, Content:Lyophilisate: 1 dose + suspension: 1 ml
Packaging:Lyophilisate: vial (glass); suspension: vial (glass), Package_size:Lyophilisate: 25 vials + suspension: 25 vials, Content:Lyophilisate: 1 dose + suspension: 1 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
7/05/2014

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
7/05/2014

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

Documents

Combined File of all Documents

English (PDF)
Published on: 7/09/2026
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