

# Versican Plus DP lyophilisate and solvent for suspension for injection for dogs

Authorised

- Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live
- Canine distemper virus, strain CDV Bio 11/A, Live

## Product identification

**Medicine name:**

Versican Plus DP lyophilisate and solvent for suspension for injection for dogs

---

**Active substance:**

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

Canine distemper virus, strain CDV Bio 11/A, Live

---

**Target species:**

Dog

---

**Route of administration:**

Subcutaneous use

---

## Product details

**Active substance and strength:**

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

19952.00 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, strain CDV Bio 11/A, Live  
1258.00 50% tissue culture infectious dose / 1.00 Dose

---

**Pharmaceutical form:**

Lyophilisate and solvent for suspension for injection

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI07AD03

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Belgium

---

**Available in:**

Belgium

---

**Package description:**

(ID2) 50 Dose; 50 millilitre(s): Box (plastics) with 50 Vial (glass) each with 1 Dose and 50 Vial (glass) each with 1 millilitre(s)

(ID1) 25 Dose; 25 millilitre(s): Box (plastics) with 25 Vial (glass) each with 1 Dose and 25 Vial (glass) each with 1 millilitre(s)

---

## 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:**

20/04/2016

---

**Manufacturing sites for batch release:**

Bioveta a.s.

---

**Responsible authority:**

Federal Agency For Medicines And Health Products

---

**Authorisation number:**

BE-V496560

---

**Date of authorisation status change:**

20/04/2016

---

**Reference member state:**

Germany

---

**Procedure number:**

DE/V/0266/001

---

**Concerned member states:**

Belgium Bulgaria Cyprus Greece Hungary Italy Lithuania Luxembourg  
Netherlands Poland Portugal Romania Spain  
United Kingdom (Northern Ireland)

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.

## Combined File of all Documents

English (PDF)

Published on: 24/05/2023

[Download](#)

## Labelling

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

2613562-paren-20251101.pdf