# Biocan Novel DHPPi, Lyophilisate and solvent 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 2b, strain CPV-2b Bio 12/B, Live
- Canine parainfluenza virus 2, strain CPiV-2-Bio 15, Live

## Product identification

**Medicine name:**
Biocan Novel DHPPi, Lyophilisate and solvent for suspension for injection
Biocan Novel DHPPi, λυοφιλοποιημένο υλικό και διαλύτης για ενέσιμο εναιώρημα για σκύλους

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

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

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

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

**Target species:**
Dog

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**

Canine distemper virus, strain CDV Bio 11/A, Live
5.10 log10 tissue culture infective dose 50 / 1.00 Dose

Canine adenovirus 2, strain CAV-2-Bio 13, Live
5.30 log10 tissue culture infective dose 50 / 1.00 Dose

Canine parvovirus 2b, strain CPV-2b Bio 12/B, Live
6.60 log10 tissue culture infective dose 50 / 1.00 Dose

Canine parainfluenza virus 2, strain CPiV-2-Bio 15, Live
5.10 log10 tissue culture infective dose 50 / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate and solvent for suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Dog**

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

QI07AD04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Cyprus

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**Package description:**

Glass Vial 50 x 1.0 Dose
Glass Vial 25 x 1.0 Dose
Glass Vial 10 x 1.0 Dose

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## 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:**
Bioveta a.s.

**Marketing authorisation date:**
12/09/2014

**Manufacturing sites for batch release:**
Bioveta a.s.

**Responsible authority:**
Ministry Of Agriculture Rural Development And Environment

**Authorisation number:**
CY00478V

**Date of authorisation status change:**
23/09/2019

**Reference member state:**
Czechia

**Procedure number:**
CZ/V/0124/001

**Concerned member states:**
Bulgaria Croatia Cyprus Estonia Hungary Latvia Lithuania Poland Romania Slovakia Slovenia

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.

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