

# Biocan DHPPi+LR Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

Authorised

- Canine distemper virus, strain CDVU 39, Live
- Canine adenovirus 2, strain CAV-2-Bio 13, Live
- Canine parvovirus, strain OP-I/81, Live
- Canine parainfluenza virus, strain CPIV-2-Bio 15, Live
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Leptospira interrogans, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Grippotyphosa, Inactivated
- Rabies virus, strain SAD Vnukovo-32, Inactivated
- Algeldrate

## Product identification

### **Medicine name:**

Biocan DHPPi+LR Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

### **Active substance:**

Canine distemper virus, strain CDVU 39, Live

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

Canine parvovirus, strain OP-I/81, Live

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

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Leptospira interrogans, serovar Canicola, Inactivated  
Leptospira interrogans, serovar Grippotyphosa, Inactivated  
Rabies virus, strain SAD Vnukovo-32, Inactivated  
Algeldrate

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**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Canine distemper virus, strain CDVU 39, Live  
Canine adenovirus 2, strain CAV-2-Bio 13, Live  
Canine parvovirus, strain OP-I/81, Live  
Canine parainfluenza virus, strain CPIV-2-Bio 15, Live  
Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated  
Leptospira interrogans, serovar Canicola, Inactivated  
Leptospira interrogans, serovar Grippotyphosa, Inactivated  
Rabies virus, strain SAD Vnukovo-32, Inactivated  
Algeldrate  
2.00 milligram(s) / 1.00 milligram(s)

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**Pharmaceutical form:**

Lyophilisate and solvent for suspension for injection

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

QI07AJ02

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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:**

Poland

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**Package description:**

Available only in [Polish](#)

Available only in [Polish](#)

Available only in [Polish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Grabikowski-Grabikowska Przedsiębiorstwo Produkcyjno-Handlowo-Uslugowe Inex Sp. j.

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**Marketing authorisation date:**

21/10/2010

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2011

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**Date of authorisation status change:**

21/10/2010

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

## Documents

### Package Leaflet

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

### Summary of Product Characteristics

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