

BIOCAN DHPPi + L PLUS

Authorised

- Canine parainfluenza virus, strain CPIV-2-Bio 15, Live
- Canine parvovirus, Live
- Leptospira interrogans, serogroup Grippotyphosa, serovar Grippotyphosa, Inactivated
- Leptospira interrogans, Serogroup Canicola, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Canine adenovirus 2, Live
- Canine distemper virus, strain CDVU 39, Live

Product identification

Medicine name:

BIOCAN DHPPi + L PLUS

Active substance:

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

Canine parvovirus, Live

Leptospira interrogans, serogroup Grippotyphosa, serovar Grippotyphosa, Inactivated

Leptospira interrogans, Serogroup Canicola, serovar Canicola, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine adenovirus 2, Live

Canine distemper virus, strain CDVU 39, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

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

3.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, Live

3.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Leptospira interrogans, serogroup Grippotyphosa, serovar Grippotyphosa, Inactivated

Leptospira interrogans, Serogroup Canicola, serovar Canicola, Inactivated

32.00 log₂ geometric mean titre / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine adenovirus 2, Live

3.50 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, strain CDVU 39, Live

3.00 log₁₀ 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AA

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

19/05/2019

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2899

Date of authorisation status change:

28/05/2023

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.

Package Leaflet and Labelling

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