File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000053308

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, type 2b, strain CPV-2b Bio 12/B, Live
- Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Product identification

Medicine name:
Biocan Novel DHPPi, Lyophilisate and solvent 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

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 50% tissue culture infectious dose / 1.00 Dose

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

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

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

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AD04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Glass Vial 10 x 1.0 Dose
Glass Vial 25 x 1.0 Dose
Glass Vial 50 x 1.0 Dose

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:
2/10/2014

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/14/0049

Date of authorisation status change:
2/10/2014

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

Combined File of all Documents

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

eu-puar-czv0124001-mr-biocan_novel_dhppi-en.pdf