# CANGLOB DHLaPPi, Injekce

Authorised

- Immunoglobulins against Canine distemper virus, Canine
- Immunoglobulins against Canine parvovirus, Canine
- Immunoglobulins against Canine parainfluenza virus, Canine
- Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine

## Product identification

**Medicine name:**
CANGLOB DHLaPPi, Injekce

**Active substance:**
Immunoglobulins against Canine distemper virus, Canine

Immunoglobulins against Canine parvovirus, Canine

Immunoglobulins against Canine parainfluenza virus, Canine

Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine

**Target species:**
Dog

**Route of administration:**
Intravenous use
Subcutaneous use
Intramuscular use

## Product details

**Active substance and strength:**

Immunoglobulins against Canine distemper virus, Canine
160.00 virus neutralising unit(s) / 1.00 Dose

Immunoglobulins against Canine parvovirus, Canine
1024.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Immunoglobulins against Canine parainfluenza virus, Canine
64.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine
160.00 virus neutralising unit(s) / 1.00 Dose

---

**Pharmaceutical form:**

Injection

---

**Withdrawal period by route of administration:**

**Intravenous use:**

- **Dog**

**Subcutaneous use:**

- **Dog**

**Intramuscular use:**

- **Dog**

---

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

QI07AM

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**
Czechia

**Package description:**
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

## 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:**
Dyntec spol. s r.o.

**Marketing authorisation date:**
16/05/1995

**Manufacturing sites for batch release:**
Dyntec spol. s r.o.

**Responsible authority:**
Institute For State Control Of Veterinary Biologicals And Medicaments

**Authorisation number:**
97/337/95-C

**Date of authorisation status change:**
2/12/2010

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

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

Labelling

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

---