# EURICAN DAPPi-LR, liofilizatas ir suspensija injekcinei suspensijai ruošti šunims

Not authorised

- Canine distemper virus, strain BA5, Live
- Canine adenovirus 2, strain DK13, Live
- Canine parvovirus, strain CAG2, Live
- Canine parainfluenza virus, strain CGF 2004/75, Live

## Product identification

**Medicine name:**
EURICAN DAPPi-LR, liofilizatas ir suspensija injekcinei suspensijai ruošti šunims

**Active substance:**
Canine distemper virus, strain BA5, Live

Canine adenovirus 2, strain DK13, Live

Canine parvovirus, strain CAG2, Live

Canine parainfluenza virus, strain CGF 2004/75, Live

**Target species:**
Dog

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Canine distemper virus, strain BA5, Live
10000.00 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, strain DK13, Live
316.23 50% cell culture infectious dose / 1.00 Dose

Canine parvovirus, strain CAG2, Live
79432.80 50% cell culture infectious dose / 1.00 Dose

Canine parainfluenza virus, strain CGF 2004/75, Live
50118.70 50% cell culture infectious dose / 1.00 Dose

---

**Pharmaceutical form:**
Lyophilisate and suspension for suspension for injection

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI07AJ06

---

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

---

**Authorisation status:**
Surrendered

---

**Authorised in:**
Lithuania

---

**Package description:**
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

---

## Additional information

**Entitlement type:**

Marketing Authorisation

**Legal basis of product authorisation:**
Legal basis not covered by Directive 2001/82/EC

**Marketing authorisation holder:**
Boehringer Ingelheim Animal Health France

**Marketing authorisation date:**
19/03/2002

**Manufacturing sites for batch release:**
Boehringer Ingelheim Animal Health France

**Responsible authority:**
State Food And Veterinary Service

**Authorisation number:**
LT/2/02/1378/001-003

**Date of authorisation status change:**
10/06/2007

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.