EURICAN DAPPi-LR, Lyofilizát a suspenze pro injekční suspenzi

Not authorised

- Canine distemper virus, Live
- Canine adenovirus 2, Live
- Canine parvovirus, Live
- Leptospira interrogans, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Canine parainfluenza virus, Live
- Rabies virus, strain G52, Inactivated

Product identification

Medicine name:
EURICAN DAPPi-LR, Lyofilizát a suspenze pro injekční suspenzi

Active substance:
Canine distemper virus, Live

Canine adenovirus 2, Live

Canine parvovirus, Live

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine parainfluenza virus, Live

Rabies virus, strain G52, Inactivated

Target species:

Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, Live
6.00 log10 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live
6.30 log10 50% cell culture infectious dose / 1.00 Dose

Canine parvovirus, Live
7.10 log10 50% cell culture infectious dose / 1.00 Dose

Leptospira interrogans, serovar Canicola, Inactivated
40.00 Protective Dose / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
40.00 Protective Dose / 1.00 Dose

Canine parainfluenza virus, Live
7.10 log10 50% cell culture infectious dose / 1.00 Dose

Rabies virus, strain G52, Inactivated
1.00 international unit(s) / 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

Authorisation status:
Surrendered

Authorised in:
Czechia

Package description:
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:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
23/04/1998

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

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

Authorisation number:
97/026/98-C

Date of authorisation status change:
3/12/2025

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.