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EURICAN DAPPi - L liofilizat in suspenzija za pripravo suspenzije za injiciranje za pse

Authorised

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

Product identification

Medicine name:
EURICAN DAPPi - L liofilizat in suspenzija za pripravo suspenzije za injiciranje za pse

Active substance:
Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Leptospira interrogans, serovar Canicola, Inactivated

Canine parainfluenza virus, Live

Canine parvovirus, Live

Canine adenovirus 2, Live

Canine distemper virus, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
1.00 European Pharmacopoeia Unit(s) / 1.00 Dose

Leptospira interrogans, serovar Canicola, Inactivated
1.00 European Pharmacopoeia Unit(s) / 1.00 Dose

Canine parainfluenza virus, Live
50118.00 log10 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, Live
79432.00 log10 50% tissue culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live
316.20 log10 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, Live
10000.00 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and suspension for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI02

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

Authorisation status:
Valid

Authorised in:
Slovenia

Package description:
Available only in Slovenian
Available only in Slovenian

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:
14/11/2021

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0125/001

Date of authorisation status change:
20/09/2004

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.