

Eurican DAPPi-LR, süstesuspensiooni lüofilisaat ja suspensioon koertele

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
- Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
- Rabies virus, strain G52, Inactivated

Product identification

Medicine name:

Eurican DAPPi-LR, süstesuspensiooni lüofilisaat ja suspensioon koertele

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

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

Rabies virus, strain G52, Inactivated

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

316.00 50% cell culture infectious dose / 1.00 dose

Canine parvovirus, strain CAG2, Live

79433.00 50% cell culture infectious dose / 1.00 dose

Canine parainfluenza virus, strain CGF 2004/75, Live

50119.00 50% cell culture infectious dose / 1.00 dose

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated

1.00 Hamster protective Dose 80% / 1.00 millilitre(s)

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

1.00 Hamster protective Dose 80% / 1.00 millilitre(s)

Rabies virus, strain G52, Inactivated

1.00 international unit(s) / 1.00 millilitre(s)

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:

Estonia

Package description:

Available only in [Estonian](#)

Available only in [Estonian](#)

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:

7/08/2003

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

State Agency Of Medicines

Authorisation number:

1147

Date of authorisation status change:

30/08/2023

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.