Eurican DAPPi-LR vakcina

Not authorised

- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
- Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
- Rabies virus, Inactivated
- Canine parvovirus, Live
- Canine parainfluenza virus, Live
- Canine adenovirus 2, Live
- Canine distemper virus, Live

Product identification

Medicine name:
Eurican DAPPi-LR vakcina

Active substance:
Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
Rabies virus, Inactivated
Canine parvovirus, Live
Canine parainfluenza virus, 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, strain 16069, Inactivated
40.00 Hamster protective Dose 80% / 1.00 unit(s)/dose

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
40.00 Hamster protective Dose 80% / 1.00 unit(s)/dose

Rabies virus, Inactivated
1.00 international unit(s) / 1.00 unit(s)/dose

Canine parvovirus, Live
4.90 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Canine parainfluenza virus, Live
4.70 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Canine adenovirus 2, Live
2.50 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Canine distemper virus, Live
4.00 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Pharmaceutical form:
Powder and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI02

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

Authorisation status:
Surrendered

Authorised in:
Hungary

Package description:

Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
17/06/1998

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
28/09/2022

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet