BIOCAN DHPPi+LR, liofilizatas ir skiediklis injekcinei suspensijai ruošti

Authorised

- Leptospira interrogans, serovar Grippotyphosa, Inactivated
- Leptospira interrogans, serovar Canicola, Inactivated
- LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED
- Rabies virus, Inactivated
- Canine parainfluenza virus, Live
- Canine parvovirus, Live
- Canine adenovirus 2, Live
- Canine distemper virus, Live

Product identification

Medicine name:
BIOCAN DHPPi+LR, liofilizatas ir skiediklis injekcinei suspensijai ruošti

Active substance:
Leptospira interrogans, serovar Grippotyphosa, Inactivated

Leptospira interrogans, serovar Canicola, Inactivated

LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED

Rabies virus, 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 Grippotyphosa, Inactivated
32.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Leptospira interrogans, serovar Canicola, Inactivated
32.00 Antibody microagglutination-lytic reaction / 1.00 Dose

LEPTOSPIRA ICTEROHAEMORRHAGIAE INACTIVATED
32.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Rabies virus, Inactivated
2.00 international unit(s) / 1.00 Dose

Canine parainfluenza virus, Live
15848.90 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, Live
316228.00 50% tissue culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live
31622.80 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, Live
31622.80 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent 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:
Valid

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:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
20/10/2009

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/09/1887/001-003

Date of authorisation status change:
2/11/2014

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

Documents

RV1887.pdf