EURICAN DAP-LMULTI LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION

Not authorised

- Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
- Canine adenovirus 2, strain DK13, Live
- Canine distemper virus, strain BA5, Live
- Canine parvovirus, strain CAG2, Live
- Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

Product identification

Medicine name:
EURICAN DAP-LMULTI LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION

Active substance:
Leptospira interrogans, serovar Canicola, strain 16070, Inactivated

Canine adenovirus 2, strain DK13, Live

Canine distemper virus, strain BA5, Live

Canine parvovirus, strain CAG2, Live

Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
1.00 Hamster protective Dose 80% / 1.00 Dose

Canine adenovirus 2, strain DK13, Live
316.23 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, strain BA5, Live
10000.00 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, strain CAG2, Live
79432.80 50% tissue culture infectious dose / 1.00 Dose

Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated
1.00 Hamster protective Dose 80% / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
1.00 Hamster protective Dose 80% / 1.00 Dose

Pharmaceutical form:
Lyophilisate and suspension for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI03

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

Authorisation status:
Surrendered

Authorised in:
Bulgaria

Package description:
Plastic box of 25 vials (glass) of lyophilisate (1 dose) and 25 vials (glass) of suspension (1 ml)
Plastic box of 50 vials (glass) of lyophilisate (1 dose) and 50 vials (glass) of suspension (1 ml)
Plastic box of 10 vials (glass) of lyophilisate (1 dose) and 10 vials (glass) of suspension (1 ml)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
19/08/2015

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2599

Date of authorisation status change:
23/07/2023

Reference member state:
France

Procedure number:

FR/V/0287/001/DC

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

Documents

eu-puar-frv0287001-mr-rpe668-en.pdf